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Environmental Testing and Certification Corp. 284 Rammar, Center Parkway P.O. Box 7806 Edison, N.J. 08810-7508 201-225-0700



PRIGINAL

June 8, 1988

USEPA Region III Central Regional Laboratory 839 Bestgate Road Annapolis, mD 21401-3099

Attn: Diane Simms

Quality Assurance Officer

Re: RI/FS - duPont Newport Site

Dear Diane:

Enclosed please find the performance and audit information requested during our teleconference of June 7, 1988. ETC Corp. is a participant in the USEPA Contract Laboratory Program's Performance Evaluation Sample and On-Site Audit Program. These are the same programs used to monitor performance for those laboratories with active sample contracts.

ETC routinely performs all of the analytical and QA/QC requirements of the CLP statements of work for organics and inorganics. ETC provides a number of reporting formats for the analytical data generated, from electronic summary reports to complete technical reports containing all raw and support data. Additionally, ETC can provide TCL Analysis, both organic and inorganic, in a full CLP deliverables package as specified in Section B of the Statement of Work.

Diane Simms USEPA Region III June 8, 1988

Page 2

I sincerely hope this information will be helpful in your task. Should you have any questions or need any additional information, please do not hesitate to contact June Baker (201-225-6741) at our Edison office.

Sincerely

John E. Farrell II

Manager, Sales Development

and Technical Support

JEF:ja attachment

pc: Gerrado Amoder, USEPA Region III

Roger T. Gresh, Woodward-Clyde

June Baker, ETC/Edison

Marilyn Bracken, ETC/Edison

7n ...

May, 1988

The following list includes the On-Site External Audits performed at the ETC-Edison facility.

| 2. 3. | 860515 860807 861209 861217 | NJDEP CA UT NJDEP | X-085 specific Haz-waste certification DOH-DW & WW certification Lab cert-A-280, DW & WW |
|----------|--------------------------------------|----------------------------|---|
| 5. | 870203 | WMI | Laboratory audit |
| | • | EPA II | IFB-TCDD |
| 7. | 870404 | EPA V | WMI sites |
| 8. | 870421 | PA | DER-DW certification |
| 9. | 870428 | NY | DOH-DW & WW certification |
| 10. | 870400 | Army Corps. | Engineers, Systems audit |
| 11. | 870415 | NJDEP | X-085 & A-280 |
| | 870819 | \mathtt{FL} | DHRS-DW & WW certification |
| 13. | 871014 | NJDEP | New CV parameters cert. |
| 14. | 871028 | MKE/RMA | Systems audit |
| | 871109 | RMA/USATHAMA | |
| 16. | 871112-14 | MMI | Laboratory audit |
| 17. | 871116-17 | WI | DW & WW certification |
| 18. | 871117-19 | EPA IV | WMI, Dupont & Ciba Geigy sites |
| 19. | 871203 | MKE | Systems & documentation |
| 20. | 880210 | NJDEP | X-195 specific |
| 21. | 880218 | Dynamac | CLP Inorganics & Organics |
| 22. | 880322 | EPA II | CLP Dioxin & Organics |
| | 880323-24 | WMI | Follow-up audit |
| 24. | 880412 | EPA IV | CLP Organics/Inorganics |
| | 880413 | NY | DOH-DW & WW certification |
| 26. | 880419 | EPA V | Support for PRP-lead Site |

PERFORMANCE EVALUATION SAMPLES

RESULTS REPORTED FROM:

- SECOND QUARTER INORGANICS (CLP)
- SECOND QUARTER ORGANICS (CLP)
- NEW JERSEY DEPARTMENT OF ENVIRONMENTAL PROTECTION
- PENNSYLVANIA DEPARTMENT OF ENVIRONMENTAL RESOURCES



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY WASHINGTON, D.C. 20460

AUG 1 | 1986

OFFICE OF
SOLID WASTE AND EMERGENCY RESPONS

Ro. 9X

'IEMORANDUM

SUBJECT: State Requirements for Laboratory Aupport

FROM:

Henry L. Longest II, Director

Office of Emergency and Remedia Response

TO:

Regional Waste Management Division Directors

Regional Environmental Services Division Directors

It has been brought to my attention that an increasing number of States are linking their requirements for Superfund analytical laboratory support to requirements of the Contract Laboratory Program (CLP). In fact, some States are apparently requiring that laboratories must be active CLP participants in order to do analytical work for the State. A number of private laboratories are concerned about such State requirements, since they believe it raises issues of equity for laboratories who are not active in the CLP for reasons other than quality, e.g. limited bid awards due to EPA funding constraints, or other factors. Obviously, those labs feel that their markets may be limited if active CLP participation is a requirement for other business.

I believe the concerns that are being raised have merit. The CLP is not intended to be a lab certification program. There are certainly laboratories outside of the CLP capable of performing high quality analytic work. However, appropriate quality assurance oversight and quality control practices, such as those applied to CLP labs, should be required for any laboratory doing Superfund work.

I suggest that you raise this issue with the States in your respective Regions and point out to them the concerns that are raised if CLP participation by laboratories is a requirement for other work.

cc: CLP Laboratories



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

OFFICE OF RESEARCH AND DEVELOPMENT ENVIRONMENTAL MONITORING SYSTEMS LABORATORY-LAS VEGAS P.O. BOX 93478 LAS VEGAS, NEVADA 89193-3478 (702/798-2100 · FTS 545-2100)

Organia... (Red) APR 1 1 1988

Diane Foster Environmental Testing & Certification Corp. 284 Raritan Center Parkway Edison, NJ 08818

Dear Ms. Foster:

For your information and review, enclosed are the results for your participation in the EMSL-LV Second Quarter Inorganic Performance Evaluation Study (QB2 FY-88). The samples were prepared by the EMSL-LV and consisted of one soil sample and two water samples. The homogeneous soil sample and one of the water samples were spiked with inorganic parameters. The other water sample was a blank. The samples were to be prepared and analyzed by current IFB procedures as per contract. All laboratories received the samples single blind. Enclosed is general information about the Superfund Performance Evaluation Program which explains the new PE portion of the Laboratory Profile Package, called the "Individual Laboratory Summary Report" (ILSR).

The EMSL-LV thanks you for your participation in this study. We trust that this information is vital to you as a member of the community of laboratories analyzing hazardous waste samples for Superfund.

Sincerely

Larry Butler, Ph.D.

Supervisor, Performance Evaluation Program
Quality Assurance Research Branch
Quality Assurance and Methods Development Division

Enclosure

cc: (w/enclosure)
Carla Dempsey, OERR
William Langley, OERR

Enclosure 1A

The EMSL-LV is adhering to the National Program Office guidelines with the following requirement. For each parameter which you failed to correctly identify or quantitate or which you reported as a false positive (parameters not added into this PE sample, but found by your laboratory at concentrations exceeding contract requirements), please document in a letter to your Project Officer, Deputy Project Officer and myself within two weeks of receipt of this letter, the source of the problem(s) and the corrective action(s) taken to prevent the problem from occurring in future quarterly blind PE samples.

Details of the new scoring procedure are shown on the following "Attachment 1." For your convenience, included here is the Individual Laboratory Summary Report (ILSR) for your laboratory and a graphical programmatic summary of scores.

C-85 . 131261

> The following information explains the details about the Individual Laboratory Summary Report, Program Summary Report, Summary of Laboratory Scores, and specific information about the scoring procedures.

The Scoring Procedures

The confidence interval (CI) calculation and the scoring algorithm are the intrinsic parts of the Quarterly Blind (QB) study. At present, the 95 percent CI are calculated from CLP laboratory-submitted results. All reported results are compared to the CI. Elements that were found to be mis-identified, misquantitated and reported false positives are flagged and used in the calculation of the score. False positives are values at exceedingly high concentrations which can be caused by contamination or interference. In addition, matrix spike accuracy and duplicate precision are included in the scoring. Other details are explained in the footnotes which accompany the Individual Laboratory Summary Report.

Confidence intervals were calculated from the laboratorysubmitted values using the statistical procedure Biweight which does not generate outliers. Instead, the laboratory-reported results are weighted relative to their position from the mean.

The following equation is used to calculate the percent score (Z score) for each laboratory.

% Score = 100 - (
$$5A + B + 2C$$
)
- ($5A^{W} + B^{W} + 2C^{W}$)
- $0.5S^{S}$ - D

where A = number of mis-identifications

$$B = \begin{bmatrix} 1 & - & T & -x \\ --- & T \end{bmatrix} * 50$$

T = total number of elements

x = number of mis-quantitations

C = number of false positives

S = number of matrix spikes outside the criteria

D = number of duplicates

outside the criteria

w = water matrix

s = soil matrix



The Scoring Procedures (continues)

The following scoring categories are recommended by the Environmental Monitoring System Laboratory, Las Vegas (EMSL-LV) under the directive of the National Program Office:

- 1. 100 to 90 percent Acceptable Performance, No Corrective Action Necessary
- 2. 90 to 75 percent Acceptable Performance, Corrective Action Necessary
- 3. below 75 percent Unacceptable Performance,
 Corrective Action Mandatory

A score below 75% results in the failure of a performance evaluation (PE) sample.

Individual Laboratory Summary Raport

| Header / Qualifier | Explanation | | | | |
|--------------------|--|--|--|--|--|
| LABORATORY NAME | laboratory name and location (state) and assigned alpha-numeric code | | | | |
| PERFORMANCE LEVEL | laboratory performance falls into one of three (3) categories: | | | | |
| | ACCEPTABLE % score greater than or equal to 90 | | | | |
| | ACCEPTABLE Z score greater - Corrective than or equal Action to 75 and less Necessary than 90 | | | | |
| | UNACCEPTABLE % score is less - Corrective than 75 Action Mandatory | | | | |
| LABORATORY RANK | comparison of CLP laboratories only for which a % score was calculated | | | | |
| · | Above number of laboratories whose %%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%%% | | | | |
| | Same number of laboratories whose % score is the equal to the laboratory's % score | | | | |
| | Below number of laboratories whose %% score is less than the laboratory's %% score | | | | |
| Z SCORE | percent score calculated using the scoring equation | | | | |
| REPORT DATE | date that the Individual Laboratory Summary report is printed and in the format, month/day/year (for example, 1/23/88) | | | | |
| MATRIX | sample matrix (water or soil) AR30398 | | | | |



Individual Laboratory Summary Report (Continued)

| Header / Qualifier | Explanation |
|--------------------|--|
| ELEMENT NAME | the 23 target analytes required by the Statement of Work |
| 95 % CI | 95 percent confidence interval (CI) calculated for each element using the Biweight procedure with CLP laboratory-submitted results |
| LOWER | lower limit of CI |
| UPPER | upper limit of CI |
| LAB RESULTS | laboratory-reported values and qualifiers |
| REPORTED VALUE | laboratory-reported concentration |
| QUALIFIER CODE | laboratory-reported qualifier(s) pertaining to the preceding value |
| PROGRAM DATA | pertains to only CLP laboratory-submitted values |
| # LABS MIS-ID | number of CLP laboratories which mis-identified the element |
| # LABS MIS-QUAN | number of CLP laboratories which mis-quantitated the element |
| # LABS FALSE POS | number of CLP laboratories which reported the element at an exceedingly high concentration |
| TOTAL # LABS | number of CLP laboratories whose values were used in the statistical study of the program data |

Individual Laboratory Summary Report (continued)

| Header / Qualifier | <u>Explanation</u> | | | | | | |
|---------------------------------|--|--|--|--|--|--|--|
| # OF ELEMENTS MIS-IDENTIFIED | number of elements mis-identified by the laboratory | | | | | | |
| # OF ELEMENTS MIS-QUANTIFIED | number of elements mis-quantitated by the laboratory | | | | | | |
| # OF FALSE POSITIVES | number of elements reported at an exceedingly high concentration by the laboratory | | | | | | |



Attachment 1 Page 6

Program Summary Report

| Header / Qualifier | Explanation |
|--------------------|--|
| MATRIX | sample matrix (water or soil) |
| REPORT DATE | <pre>date that the Program Summary Report is printed and in the format, month/day/year (for example, 1/23/88)</pre> |
| ELEMENT DATA | element data generated with CLP laboratory-submitted results |
| ELEMENT NAME | the 23 elements required by the Statement of Work |
| SPIKE LEVEL | the level spiked into the sample |
| 95 % CI | 95 percent confidence interval (CI) calculated for each element using the Biweight procedure with CLP laboratory-submitted results |
| LOWER | lower limit of CI |
| UPPER | upper limit of CI |
| MEAN RESULT | average/mean of the values used in the calculation of the CI |
| STANDARD DEVIATION | standard deviation of the values used in the calculation of the CI |
| PROGRAM DATA | pertains to only CLP laboratory-submitted values |
| # LABS MIS-ID | number of CLP laboratories which mis-identified the element |
| # LABS MIS-QUAN | number of CLP laboratories which mis-quantitated the element |
| # LABS FALSE POS | number of CLP laboratories which reported the element at an exceedingly high concentration |
| TOTAL # LABS | number of CLP laboratories whose |

values were used in the statistical study of the program data

Attachment 1 Page 7

Program Summary Report (continues)

Header / Qualifier

OF LABS WITH ACCEPTABLE PERFORMANCE

OF LABS WITH ACCEPTABLE PERFORMANCE CORRECTIVE ACTION NECESSARY

OF LABS WITH UNACCEPTABLE PERFORMANCE -CORRECTIVE ACTION MANDATORY

Explanation

number of CLP laboratories whose % score is greater than or equal to 90

number of CLP laboratories whose % score is less than 75

Attachment 1 Page 8



Summary of Laboratory Scores

| Header / Qualifier | <u>Explanation</u> |
|--------------------|--|
| LAB NAME | SMO assigned laboratory lab code |
| CODE | assigned alpha-numeric laboratory code |
| SCORE | % score calculated for each laboratory |
| MIS-ID | number of elements mis-identified (the "A" in the % Score equation) |
| MIS-QUANT | number of elements mis-quantified (the "B" in the % Score equation) |
| FALSE POS | number of false positives reported (the "C" in the % Score equation) |
| MSPK OUT | number of matrix spike recoveries outside the criteria (the "S" in the % Score equation) |
| DUP OUT | number of duplicates (RPDs) outside the criteria (the "D" in the % Score equation) |

(Red)

INORGANIC PERFORMANCE EVALUATION SAMPLE INDIVIDUAL LABORATORY SUMMARY REPORT FOR QB 2 FY 8B

LABORATORY NAME: Env. Testing & Certif. (NJ) [P2]

PERFORMANCE LEVEL: ACCEPTABLE

LABORATORY RANK: Above = 5 Same = 1 Below = 24

% Score: 96.6 REPORT DATE: 3/23/1988

MATRIX: SOIL

| | | | LAB F | RESULTS | | | PROGR | AM DATA | | |
|--------------|-------|------------------|-------------------|-----------|--------|---------------|-----------|--------------|---------|-------|
| ELEMENT NAME | 95 | 1 CI | REPORTED | QUALIFIER | *LABS | \$LABS | #LABS | ILABS | *LABS | TOTAL |
| | LOWER | UPPER | VALUE | CODE | MIS ID | MIS-QUANT | FALSE POS | MSPK OUT | DUP OUT | *LABS |
| ALUHINUM | 4798 | 11900 | 9668 | | • | 2 | • | • | • | 31 |
| ARTIMONY | 8 | 53 | 23 | | 3 | 3 | • | 20 | • | 31 |
| ARSENIC | 17 | 28 | 25 | | • | 4 | 0 | 7 | 1 | 31 |
| BARIUH | 156 | 189 | 179 | | • | 3 | 0 | 1 | • | 31 |
| BERYLLIUM | 16 | 21 | 18 | | • | • | • | 1 | 0 | 31 |
| CADMIUM | 9.7 | 17 | 13 | | • | • | 8 | 1 | 0 | 31 |
| CALCIUM | 75301 | 194991 | 93000 | | 0 | 2 | • | 9 | • | 31 |
| CHRONIUM | 16 | 51 | 42 | | • | 2 | 8 | 8 | 8 | 31 |
| COBALT | 71 | 92 | 79 | | 9 | 1 | 0 | 8 | 8 | 31 |
| COPPER | 88 | 112 | 99 | | 8 | 3 | 6 | 1 | 8 | 31 |
| IRON | 12600 | 17400 | 17200 | | • | 3 | 8 | • | • | 31 |
| LEAD | 164 | 226 | 186 | | • | 4 | • | 2 | 0 | 31 |
| MAGNESIUM | 40801 | 57101 | 529 00 | | 9 | 2 | 0 | 6 | 0 | |
| MANGARESE | 2810 | 3530 | 3570 | X | • | 7 | 8 | 1 | 8 | |
| HERCURY | 12 | 24 | 15 | | • | 3 | • | 2 | 1 | 31 |
| NICKEL | 26 | 54 | 43 | | • | 2 | • | 3 | 0 | 31 |
| POTASSIUM | 8 | 197 0 | 158 6 | | • | 4 | • | • | • | 31 |
| SELENIUM | 6.5 | 20 | 14 | | • | 3 | 9 | 4 | 4 | 31 |
| SILVER | 33 | 52 | 46 | | • | 3 | • | 5 | 1 | 31 |
| SODIUM | d | đ | 282 | | • | • | • | • | • | 31 |
| THALLIUM | 19 | 43 | 31 | | • | • | • | 6 | 2 | 31 |
| VANADIUH | 41 | 78 | 59 | | • | 1 | • | • | • | 31 |
| ZIMC | 162 | 289 | 189 | | • | 2 | • | 2 | • | 31 |

* OF ELEMENTS NOT IDENTIFIED: *

OF ELEMENTS MISQUANTIFIED: 1

• OF FALSE POSITIVES: •

• OF DUPLICATES OUT: •

WATER : SOIL :

OF MATRIX SPIKES OUT: 0

WATER : SOIL :



INORGANIC PERFORMANCE EVALUATION SAMPLE INDIVIDUAL LABORATORY SUMMARY REPORT FOR OB 2 FY 88

LABORATORY NAME: Env. Testing & Certif. (NJ) [P2]

PERFORMANCE LEVEL: ACCEPTABLE

LABORATORY RANK: Above = 5 Same = 1 Below = 24

1 Score: 96.6 REPORT DATE: 3/23/1988

MATRIX: WATER

| | | | LAB R | ESULTS | | | PROGR | AH DATA | | |
|--------------|-------|-------|--------------------|--------|--------|--------------|-----------|----------|---------|-------|
| ELEMENT MAME | 95 | Z CI | REPORTED QUALIFIER | | *LABS | !LABS | *LABS | •LABS | *LABS | TOTAL |
| | LOVER | UPPER | VALUE | CODE | HIS ID | HIS-QUANT | FALSE POS | MSPK OUT | DUP OUT | PLABS |
| ALUHINUN | 2540 | 3388 | 2888 | | • | 1 | • | 0 | 0 | 31 |
| Antimony | 4 | 111 | 91 | | 3 | • | 8 | 1 | 3 | 31 |
| ARSENIC | 68 | 106 | 88 | | 8 | 1 | • | 8 | 8 | 31 |
| BARIUM | 372 | 450 | 407 | | • | 4 | 8 | 6 | 1 | 31 |
| BERYLLIUM | 38 | 51 | 43 | | • | 1 | • | • | 8 | 31 |
| CADMIUM | 19 | 32 | 26 | | 8 | • | • | 8 | 1 | 31 |
| CALCIUM | 12300 | 15500 | 13500 | | • | 2 | • | • | 8 | 31 |
| CHRONIUM | 14 | 40 | 27 | | 8 | • | • | 8 | 1 | 31 |
| COBALT | 66 | 113 | 96 | | • | • | • | . 9 | 0 | 31 |
| COPPER | 189 | 244 | 204 | | 0 | 2 | 0 | 1 | 2 | 31 |
| IPON | 355 | 442 | 396 | | 8 | 4 | • | 0 | 0 | 31 |
| LEAD | 12 | 25 | 16 | | • | • | 6 | 3 | 2 | 31 |
| HACHESIUM | 7836 | 9600 | 8610 | | • | 2 | 8 | 6 | 0 | 31 |
| NESE | 62 | 81 | 68 | | • | 1 | • | 9 | 8 | 31 |
| URY | 10 | 28 | 16 | | • | 2 | • | 1 | 1 | 31 |
| NICKEL | 86 | 126 | 103 | | • | 1 | 8 | 0 | 1 | 31 |
| POTASSIUM | 8810 | 12400 | 18200 | | • | 2 | • | • | 8 | 31 |
| SELENIUN | 18 | 28 | 24 | | 8 | 2 | 8 | 1 | • | 31 |
| SILVER | C | c | 7.2 | | • | • | • | 5 | • | 31 |
| SODIUN | 6100 | 8320 | 6998 | | . • | 5 | • | 6 | • | 31 |
| THALLIUM | 51 | 88 | 68 | | • | 1 | • | 7 | 1 | 31 |
| VANADIUN | 118 | 154 | 135 | | • | 1 | • | 1 | • | 31 |
| ZINC | 47 | 66 | 58 | | • | 5 | • | 1 . | 2 | 31 |

* OF ELEMENTS NOT IDENTIFIED: 8

* OF ELEMENTS MISQUANTIFIED: 0

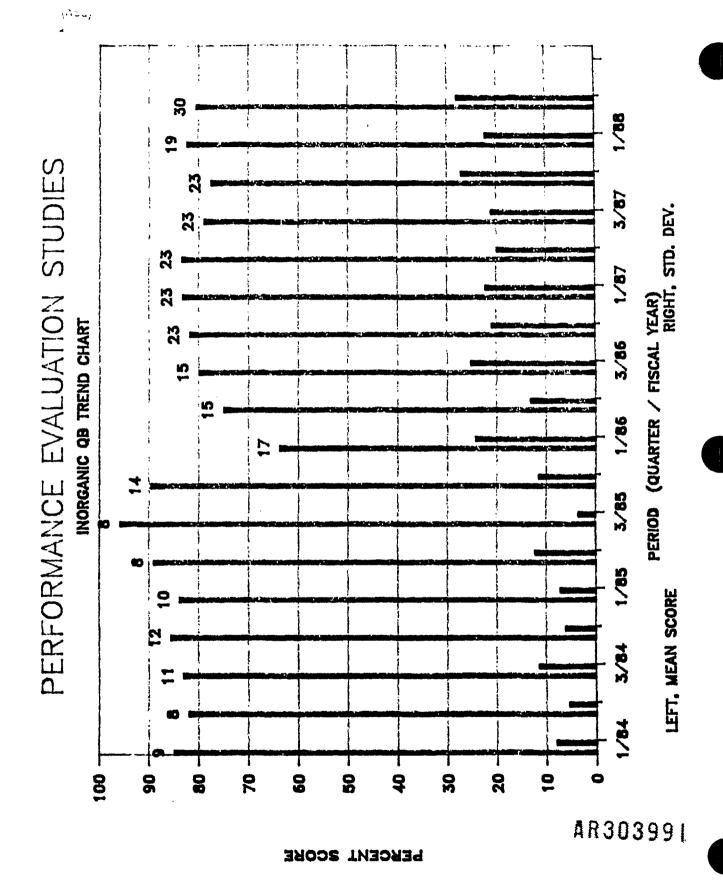
OF FALSE POSITIVES: 0

OF DUPLICATES OUT: 0

WATER : SOIL :

OF MATRIX SPIKES OUT:

WATER :



CA 2 FT 88 INORCANIC, CASE NO. 8782

D CONTIDENCE INTERVALE (CI) WERE DERIVED FROM LABORATORY SUBMITTED VALUES.

9-VALUES, AND NON-SUBMITTED VALUES (-) WERE NOT USED IN THE CALCULATION OF THE CI.

C I FERE DOT SEE STORME NOTES, PROCEDURE FOR CALLUES NO. 4.

6 CI NOT USED. SEE SCORING NOTES, PROCEDURE FOR CALLUES NO. 4.

1 INDICATES AN ESTIMATED VALUE LESS THAN THE CED. SHE AS II-FLAG.

1 INDICATES A PALLE ESTIMATED WIND ELSS THAN THE CED. SHE AS II-FLAG.

1 INDICATES A VALUE DETERMINED BY THE WETWOO OF STANDARD ADDITION.

1 INDICATES VALUE DETERMINED BY THE WETWOO OF STANDARD ADDITION.

1 INDICATES VALUE DETERMINED BY THE WETWOO OF STANDARD PROCEDD.

1 VALUE NAS OFFSIDE THE VARIENC LINIT ONLY. NO POINTS DEDUCTED.

1 VALUE NAS OFFSIDE THE VARIENC LINIT ONLY. NO POINTS DEDUCTED.

1 INDICATES A FALSE POSITIVE BY DIDAYS TEST. POINTS DEDUCTED.

1 INDICATES A RAISE DESTINATED VALUE LESS THAN THE CEDL. SAME AS B-FLAG.

1 INDICATES AN ESTIMATED VALUE LESS THAN THE CEDL. SAME AS B-FLAG.

1 INDICATES AN ESTIMATED VALUE LESS THAN THE CEDL. SAME AS B-FLAG.

1 INDICATES AN ESTIMATED VALUE LESS THAN THE CEDL. SAME AS B-FLAG.

1 INDICATES AND SULFIED BY A FACTOR OF 10.

1 INDICATES AND SULFIED BY A FACTOR OF 10.

2 THE SAMPLE MAS DILUTED BY A FACTOR OF 10.

2 THE SAMPLE MAS DILUTED BY A FACTOR OF 40.

3 AND THE SAMPLE WAS DILUTED BY A FACTOR OF 40.

3 AND THE SAMPLE WAS DILUTED BY A FACTOR OF 40.

4 AND THE SAMPLE WAS DILUTED BY A FACTOR OF 50.

5 AND THE SAMPLE WAS DILUTED BY A FACTOR OF 50.

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5 AND THE SAMPLE WAS DILUTED BY A FACTOR OF 50.

5 AND THE SAMPLE WAS DILUTED BY A FACTOR OF 50.

PROCEDURE FOR CRADING N-VALVES

ANY W-VALUE RESPONSE (INSTRUMENT DETECTION LIMIT) > CROL FOR THE APPROPRIATE DILUTION, EVEN IF IT IS IN THE 95 I CI, CAUSES A POINT DEDUCTION. IF 25 I OR MORE OF THE LABORATORIES REPORT A U-VALUE OVER THE CROL, NO POINTS ARE DEDUCTED FOR ANY LABORATORY, POSSIBLY INDICATING A MATRIX INTERFERENCE IN THE SAMPLE.

IF CHOL < LOWER CI, THEN WSE CI AS SET.

~;

IF LOWER CI < CIDE AND CIDE - UPPER CI, THEN SET LOWER CI TO ZERO (0). NO POINTS DEDUCTED FOR IDENTIFICATION OR QUARTITATION LESS THAN OR EQUAL TO THE CROS. ä

IF CRDI. > LOWER AND WPPER CI, THEN NO CI USED. PARAMETER DROPPED FROM THE SCORING. NO POINTS DEDUCTED FOR IDENTIFICATIONS OR QUANTITATIONS. FALSE POSITIVES POSSIBLE.

NOTE THAT ONLY CLP LABORATORIES WERE USED IN THE CALCULATION OF THE CI.

NOTE THAT A 9-VALUE FOLLOWED BY X (U X) HEARS THAT POINTS WERE LOST FOR IDENTIFICATION AND QUANTITATION.



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

RECEIVED APP 2 9 1929 OFFICE OF RESEARCH AND DEVELOPMENT ENVIRONMENTAL MONITORING SYSTEMS LABORATORY-LAS VEGAS P.O. BOX 93478

LAS VEGAS. NEVADA 89193-3478 (702/798-2100 - FTS 545-2100)

Mr. Jack Farrell Environmental Testing and Certif. Corporation 284 Raritan Center Parkway Edison, NJ 08818

(Red)

Dear Mr. Ferrell:

For your information and review the results for your participation in the EMSL-LV Second Quarter Organic Performance Evaluation Study (QB2, FY 88) are included here. Enclosed is general information about the Superfund Performance Evaluation Program. The PE portion of the Laboratory Profile Package, called the "Individual Laboratory Summary Report" (ILSR) was described in your letter reports last quarter. Other general information about the PE program is explained on the following pages.

The samples consisted of aqueous materials spiked with Target Compound List (TCL) and non-TCL pollutants at environmentally representative levels. Samples for all laboratories were from the same homogeneous batch. Each sample set was to be prepared and analyzed by current contractually required procedures.

The EMSL-LV thanks you for your participation in this study and wishes to congratulate the laboratories for an overall fine performance. We trust that this information is vital to you as a member of the community of laboratories analyzing hazardous waste samples for Superfund.

Sincere1

Larry Butler, Ph.D.

Supervisor, Performance Evaluation Program Quality Assurance Research Branch

Quality Assurance and Methods Development Division

Enclosure

cc: (w/enclosure) Carla Dempsey, OERR Joan Fisk, OERR Emile Boulos, OERR Angelo Carasea, OERR Howard Fribush, OERR

Enclosure

The sample set consisted of aqueous materials spiked with base/neutral/acid/pesticide (BNAP) Target Compound List (TCL) and non-TCL compounds diluted in water to environmentally representative levels (full-volume organics). This included three (3) 80-ounce bottles of semi-volatiles and pesticides; one (1) 80-ounce bottle filled with blank water for BNAP blank analyses; four (4) 40-mL vials filled with water spiked with volatile organics; and two (2) 40-mL vials filled with blank water for volatiles blank analysis. The sample set was to be prepared and analyzed by current contractually required procedures.

All analytical results, calibrations, quality control procedures, and reporting and deliverable requirements were to be submitted by the participating laboratories by contract as a regular case.

EMSL-LV PE Reports - The entire format for EMSL-LV PE reports has been revised. Identification, Quantification, and Contamination (formerly called false positives) are now scored by by an algorithm contained in your laboratory's "Individual Laboratory Summary Report" (ILSR).

Confidence Intervals (CI) were derived from the laboratory submitted values using the statistical procedure BIWEIGHT which does not generate outliers. Instead values are weighted as to their position, relative to the mean. No values are discarded. Other details are included in your ILSR. The confidence interval calculation and the scoring algorithm are intrinsic parts of the ILSRs.

Also in the footnotes to the study is the EMSL-LV method for the scoring of U-flagged values. This U-value scoring procedure has not changed from earlier PE studies.

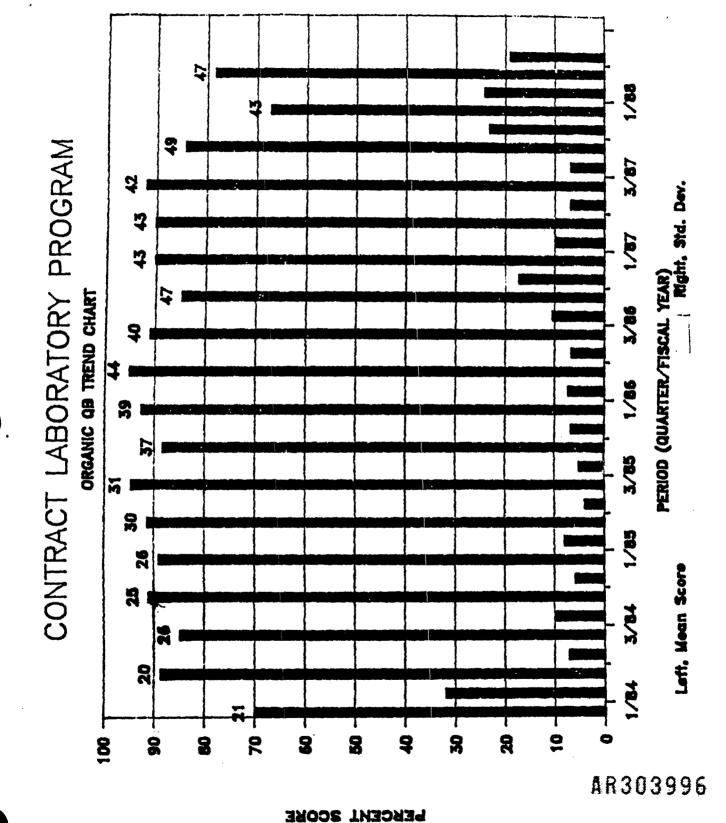
For your convenience, attached are the ILSR for your laboratory, footnotes, and a graphical programmatic representation of scores. The bar graph shows the mean laboratory performance plotted versus time. The left bar for each quarter represents the mean score, whereas the right bar for the same quarter is the standard deviation of the scores. The numbers on top of the left bar are the numbers of laboratories in each study. Please compare your score with the programmatic mean.

The EMSL-SK is recommending the following scoring categories, which are a National Program Office directive:

- 1. 100 to 90 percent "Acceptable Performance, No corrective action necessary;"
- 2. 90 to 70 percent "Acceptable Performance,
 Corrective Action Necessary;"
- 3. 70 percent or lower "Unacceptable Performance,
 Corrective Action Mandatory."

The Analytical Operations Branch of the Office of Emergency and Remedial Response also requires that all laboratories who fail to correctly identify or quantify two or more parameters or compounds or who have blank contamination (false positives) exceeding the contract requirements document the corrective action they plan to undertake. These laboratories must document in a letter to their Project Officer, Deputy Project Officer, and myself within two weeks of receipt of the results of this study, the source of the problem(s) and the corrective action(s) the laboratory plans to implement to prevent the problem(s) from occurring in future Quarterly Blind PE samples.

The government reserves the right to fairly and equably adjust scores for any PE study, should the National Program Office determine that there were unusual problems with the PE samples themselves or the scoring procedure. Determinations made by the National Program Office are final.



ORGANIC PERFORMANCE EVALUATION SAMPLE INDIVIDUAL LABORATORY SUMMARY REPORT FOR QB 2 FY 88

LABORATORY: Env. Testing & Certif. (NJ)
PERFORMANCE: ACCEPTABLE - Corrective Actions Necessary
RANK: Above = 13 Same = 2 Below = 35

* SCORE: 67.3 REPURI DATE: 4 1/1966 MATRIX: WATER

| , | | | | MAEU . | | | | |
|---|--|--|--|-------------|---|----------------------------------|---|--|
| СОМРОИМО | 90 Lower | # CI UPPER | LABORA' DAT. CONC | | #LABS NOT-ID | PROGRAM ≉LABS HIS-QUANT | 4 DATA ≉LAB3 CUNTAH | TOTAL *LABS |
| TCL VOLATILE | | | | | | | | |
| BROMCHETHANE MEIHYLENE CHLORIDE 1,1-DICHLOROETHANE 2-BUTANONE BROMODICHLOROMETHANE 1,1,2-TRICHLOROETHANE BENZENE 2-HEXANONE TOLUENE CHLOROBENZENE STYRENE XYLENES (TOTAL) | 64 c 34 38 59 54 12 48 18 85 80 129 | 240 c 555 170 80 76 17 200 30 110 110 180 | 110 120 48 110 64 62 146 89 84 | В | 000300110000 | 20 37 38 53 23 65 | 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 | 55555 55555555555555555555555555555555 |
| TCL SEMIVOLATILE | | | | | | | | |
| 2-CHLOROPHENOL N-NITROSO-DI-N-PROPYLAMINE ISOPHORONE 2.4-DIMETHYLPHENOL BENZOIC ACID HEXACHLOROBUTADIENE 2-METHYLNAPHTHALENE 2.4.6-TRICHLOROPHENOL 2-HITROANILINE ACENAPHTHYLENE ACENAPHTHYLENE ACENAPHTHENE 2.4-DINITROPHENOL DIBENZOFURAN 4-HITROPHENOL FLUORENE DIETHYLPHTHALATE PENTACHLOROPHENOL PHENANTHRENE ANTHRACENE PYRENE BUTYL BENZYL PHTHALATE BENZO(A)ANTHRACENE DI-N-OCTYL PHTHALATE DIBENZ(A,H)ANTHRACENE | 23 45 165 160 161 17 181 181 181 181 181 181 181 181 181 | 52 84 140 53 200 160 555 100 100 100 260 160 200 100 100 100 100 100 100 100 100 10 | 36 63 100 28 40 110 36 84 78 81 83 160 150 91 84 26 210 86 89 91 92 82 200 | J J | 666666666666666666666666666666666666666 | 5652723828476149654682222 | 000000000000000000000000000000000000000 | 5000000000000000000000000000000000000 |
| TCL PESTICIDES | | | | <i>)</i> | | | | |
| HEPTACHLOR ALDRIN ENDRIN TOXAPHENE | 0.05 0.13 0.16 c | 6.43 6.53 6.48 c | 0.15 0.38 0.56 5.9 | CX | 19 3 | 8 5 11 9 | 8 8 1 | 50 50 50 50 |
| NON-TCL SEMIVOLATILE | | | | | | | | |
| BENZOPHENONE DISULFOTON CHLORPYRIFOS 2-NITRO-P-CRESOL | | | 158 48 38 58 | J J J | 8 8 0 | ÅR3 | 03997 | 50 50 50 50 |
| TCL VOLATILE (Contaminants) | | | | | | | | |
| ACETONE | | | 49 | В | 0 | 9 | 8 | 50 |



ORGANIC PERFORMANCE EVALUATION SAMPLE INDIVIDUAL LABORATORY SUMMARY REPORT FOR QB 2 FY 88

* SCCRE: 87.3 REPORT DATE: 4/1/1998 MATRIX: WATER

| COMPOUND | 99 % CI LOWER UPPER | } } | LABORAT DATA CONC | | | *LABS NOT-ID | PROGRAM #LABS MIS-GUANT | DATA #LAFS CONTAM | TOTAL #LAco |
|--------------------------------------|------------------------|--------|----------------------------|----------------|------|-----------------|-------------------------------|-------------------------|----------------|
| TCL SEMIVOLATILE (Contaminants) | | | | | | | | | |
| BENZYL ALCOHOL | | | 8 | j | | 8 | 8 | ð | 58 |
| NON-TCL VOLATILE (Contaminants) | | | | | | | , | | |
| HEXANE | | | 10.2 | JB | | 9 | 8 | 6 | 5ê |
| NON-TCL SEMIVOLATILE (Contaminants) | | | | | | | : | | |
| unknown Unknown Unknown | | | 3 0 23 22 | BJ JF JF | `, | 9 9 | 0 0 | 18 18 4 | 50 50 50 |
| * OF TCL COMPOUNDS NOT-IDENTIFIED: 0 | | _ | <u></u> | | | • | | | |

^{*} OF TCL COMPOUNDS MIS-QUANTIFIED: 2 * OF TCL CONTAMINANTS: 0

 $[\]pmb{*}$ OF NON-TCL COMPOUNDS NOT-IDENTIFIED: $\pmb{\theta}$ $\pmb{*}$ OF NON-TCL CONTAMINANTS: 2

QB 2 FY 88 ORGANIC, CASE NOS. 8783 AND 8784

TCL:

- b CONFIDENCE INTERVALS (CI) WERE DERIVED FROM LABORATORY SUBMITTED VALUES. LESS THAN VALUES (<x), J-VALUES. U-VALUES, B-VALUES, AND HON-SUBMITTED VALUES (-) WERE NOT USED IN THE CALCULATION OF THE CI. CI WERE NOT SET SINCE 40 % OR MORE OF THE LABORATORIES SUBMITTED A NON-USABLE VALUE. INDICATES THAT THE COMPOUND WAS FOUND IN THE BLANK.

- INDICATES A DILUTION.
 COMPOUND EXCEEDS CALIBRATION RANGE OF INSTRUMENT.
 ESTIMATED VALUE LESS THAN THE CROL.
 NOT APPLICABLE OR NOT ANALYZED FOR.
 NOT REQUIRED.
 NOT SUBMITTED.

- ANALYZED FOR BUT NOT DETECTED.
- VALUE WAS OUTSIDE BOTH THE WARNING AND THE ACTION LIMIT. POINTS DEDUCTED FOR QUANTITATION ONLY, POINTS DEDUCTED FOR IDENTIFICATION ONLY.

- VALUE WAS OUTSIDE THE WARNING LIMIT ONLY. NO POINTS DEDUCTED.
 VALUE NOT SUBMITTED FOR THIS COMPOUND.
 INDICATES A TCL CONTAMINANT DETERMINED BY GRUBB'S TEST FOR COMPOUNDS WITH NO CI SET BASED ON "c" CRITERIA.
 BEST ESTIMATE OF VALUE AND/OR QUALIFIER. POOR OR ILLEGIBLE COPY SUBMITTED.
 WARNING LIMIT (80 PERCENT CI).
- ## ACTION LINIT (90 PERCENT CI).

HON-TCL / TIC:

- NA NOT APPLICABLE. POINTS WERE NOT DEDUCTED SINCE 40 PERCENT OF THE LABORATORIES DID NOT IDENTIFY THIS COMPOUND. NOT IDENTIFIED.
- NO NOT DETECTED. POINTS DEDUCTED.

'

- INDICATES A CONTANINANT. POINTS DEDUCTED. INDICATES THAT THE DATA WERE MANUALLY MARIPULATED BY THE ANALYST.
- A ALDOL CONDENSATION PRODUCT.

PROCEDURE FOR GRADING U-VALUES SCORING NOTES:

- 1. ANY U-VALUE RESPONSE (LABORATORY DETECTION LIMIT) > CROL, EVEN IF IT IS IN THE 90 % CI, CAUSES A POINT DEDUCTION. IF 25 % OR MORE OF THE LABORATORIES REPORT A U-VALUE OVER THE CROL, THEN MO POINTS ARE DEDUCTED FOR ANY LABORATORY. THIS COULD INDICATE A MATRIX INTERFERENCE IN THE SAMPLE.
- 2. IF CROL < LOWER CI, THEN USE CI AS SET.
- 3. IF LOWER CI < CROL AND CROL < UPPER CI, THEN SET LOWER CI TO ZERO (0). NO POINTS DEDUCTED FOR IDENTIF ATION OR QUANTITATION LESS THAN OR EQUAL TO THE CRDL.
- 4. IF CROL > LOWER AND UPPER CI, THEN NO CI USED. ANALYTE DROPPED FROM THE SCORING. NO POINTS DEDUCTED FOR IDENTIFICATIONS OF QUANTITATIONS. CONTAMINANTS POSSIBLE.

NOTE THAT ONLY CLP LABORATORIES WERE USED IN THE CALCULATION OF THE CI.

NOTE THAT A U-VALUE FOLLOWED BY AN AMPERSAND (4) (U 4) HEARS THAT POINTS WERE LOST FOR IDENTIFICATION ONLY.

NOTE THAT FOR NON-TCL/TIC A DASH FOLLOWED BY A "ND" (- ND) INDICATES THAT POINTS WERE DEDUCTED FOR IDENTIFICATION ONLY.





STATE OF NEW JERSEY

DEPARTMENT OF ENVIRONMENTAL PROTECTION

Certifies That
ENVIRONMENTAL TESTING & CERTIFICATION CORPORATION
284 Raritan Center Parkuay
Edison, NJ 08837



hating duly not the coquirements of the

Regulations Governing Laboratory Certification
And Standards Of Performance NJA.C. 7:18 et. seq.

is hereby approved as a

State Certified Water Laboratory

To perform the analyses as indicated on the Annual Certified Parameter List which must accompany this certificate to be suits



DEPARTMENT OF ENVIONILENT POSTECTION

FEMILANENT CERTIFICATION MUMBER
July 8, 1982
DATE

N.J.A.C. 7:18-2.11(d) and agreed to by the Laboratory Manager on Illing the application This certification is aubject to unannounced laboratory inspections as specified by

TO BE CONSPICUOUSLY DISPLAYED AT THE LABORATORY WITH THE ANNUAL CERTIFIED PARAMETER LIST.

7





State of Rew Jersey

DEPARTMENT OF ENVIRONMENTAL PROTECTIONDIVISION OF FINANCIAL MANAGEMENT, PLANNING & GENERAL SERVICES CN 402

TRENTON, N.J. 08625

January 21, 1988

Environmental Testing & Certification Corp. 284 Raritan Center Parkway Edison, NJ 08837

Lab ID# 12257

Dear Dr. Fitzgerald:

Enclosed is your 1987-88 Annual Certified Parameter List. This list replaces the 1986-87 form and must be conspicuously displayed with the permanent certificate at the laboratory.

Your cooperation in this matter is appreciated.

Sincerely,

Maria Salamandra, Chief

Bureau of Collections, Licensing

and Management Services

MS/DP/ch

Enclosure

cc: Jerry Bundy

STATE OF NEW JERSEY DEPARTMENT OF ENVIRONMENTAL PROTECTION OFFICE OF DUALITY ASSURANCE ANNUAL CERTIFIED PARAMETER LIST FOR 1937-1933



10 a 10

ENVIRONMENTAL TESTING AND (12257) IS CERTIFIED TO REPRORM THE ANALYSES CERTIFICATION CORP. FELOA UNITEL UNNE 30 1988.

DRINKING WATER LARDGATORY CERTIFICATION_ .

LIMITED CHEMISTRY

- 934 NITRATE, AUTO CO REDUC
- PRI FLUCRICE-AUTO ALIZ FL BLU
- A44 TURBIDITY
- 971 PH. GLASS ELECTRODE
- 956 SULFATE, GRAVIM OR TURBIO

ATOMIC ABSORPTION

- 901 34, ATOMIC ABSORPTION
- 902 AS. ATOMIC ABSORPTION
- 903 CU, ATOMIC ABSORPTION
- 904 FE. ATOMIC ABSORPTION
- 906 ZN, ATOMIC ABSORPTION
- 912 HG, MANUAL COLD VAPOR
- 914 AS. GRAPHITE FURNACE
- 915 BA+ GRAPHITE FURNACE
- 916 CD. GRAPHITE FURNACE
- 917 CR. GRAPHITE FURNACE

PAGE 1

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ATOMIC ARSORPTION

- 913 PB, GRAPHITE FURNACE
- PRO SE, GRAPHITE FURNACE
- 424 MY. GRAPHITE FURNACE
- 934 NA. ATOMIC ABSORPTION

CAS CHRIMATIGRAPHY

- 601 PURGEABLE HALDCARBONS
 - 298→ TRICHLORDETHENE
 - 2937 TETRACHLORGETHENS
 - 2982 CARBON TETRACHLORIDE
 - 2931 1,1,1-TRICHLORDETHANE
 - 2981 1,2,-DICHLORGETHANE
 - 2975 VINYL CHLORIDE
 - 2954 METHYLENE CHLORIDE
 - 2977 1,1-DICHLORDETHENE
 - 2382 TRANS-1,2-DICHLORDETHENS
 - 2989 CHLOROSENZENE
 - 2406 1.2-DICHLOROBENZENE
 - 2402 1.3-DICHLOROBENZENE
 - 2404 1,4-DICHLOROBENZENE
- 502 PURGEABLE AROMATICS
 - 2990 BENZENE
 - 2989 CHLOROBENZENE
 - 240A 1.2-DICHLOROBENZENE
 - 2402 1.3-DICHLOROBENZENE
 - 24C4 1,4-DICHLOROBENZENE
 - 2384 ORTHO-XYLENE
 - 2995 META-XYLENE
 - 2336 PARA-XYLENE
- 608 ORGANOCHLORINE PEST & PCB
 - 2959 CHLORDANE
 - 2383 AROCHLOR 1016
 - 2390 AROCHLOR 1221
 - 2392 AROCHLOR 1232
 - 2394 AROCHLOR 1242
 - 2396 AROCHLOR 1248
 - 2398 AROCHLOR 1254
 - 2400 AROCHLOR 1260

SAS CHROMATOGRAPHY

```
524 PURGEABLES
         2984 TRICHLIROETHENE
MAL
         2987 TETRACHLORDETHENE
· . . ,
         2982 CAPBON TETRACHLORIDE
         2981 1,1,1-TRICHLORDETHANE
         2930 1.2-DICHLORGETHANE
         2975 VINYL CHUDRIDE
         2964 METHYLENE CHLORIDE
         1977 1,1-DICHLORDETHENE
         2332 TRANS-1.2-DICHLORDETHENE
        2990 BENZENE
         2989 CHLDROBENZENE
 525 BASE/MEUTRALS AND ACIDS
         2406 1.2-DICHLORGBENZENE
         2402 1.3-DICHLOROBENZENE
         240+ 1.4-DICHLOROBENZENE
         2333 AROCHLOR 1016
         2390 APRCHLOR 1221
         2392 ARCCHUDR 1232
         2394 ARDCHLOR 1242
        2396 AROCHLOR 1248
        2393 AROCHLOR 1254
         2400 ARDCHLOR 1260
         2959 CHLORDANE
         2373 1,2,4-TRICHLOROBENZENS
         2384 DRITHO-XYLENE
        2995 META-XYLENE
         2386 PARA-XYLENE
      OPGANOCHLORINE PESTICIDES
 941
        VISCUE
        LINDANE
         METHOXYCHLOR
         TOXAPHENE
 942 CHLOROPHENOXY ACID HERB
         2,4-0
         2.4.5-TP(SILVEX)
 743
      TRIHALOMETHANES
         CHLOROFORM
         BROMOFORM
         BROMODICHLOROMETHANE
         DIBROMOCHLOROMETHANE
```

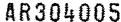
WATER POLLUTION LABORATORY CERTIFICATION

LIMITED CHEMISTRY

00076 TURBICITY

PAGE 3

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LIMITED CHEMISTRY

00095 SPECIFIC CONDUCTANCE

19349 090

20400 HYDROGEN ION-PH

70410 ALKALINITY

hospo for Jourds

20530 SUSP SOLIDS

CC556 DIL AND GREASE

00615 NITRITE

00630 NITRATE

00680 ORGANIC CARBON, TOTAL

CO720 CYANIDE, TOTAL

CC940 CHLORIDE

00945 SULFATE

00951 FLUORIDE, TOTAL

32730 PHENDLS

ATOMIC ABSORPTION

00915 CA, DISS

00916 CA, TOTAL

00925 MG, DISS

00927 MG, MITTAL

CC929 NA, TOTAL

PAGE

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AR304006

PIGNA,

ATOMIC ABSORPTION

ose ofbest. NA. DISS

> 00935 K, DISS

00937 K. TOTAL

01000 AS. DISS

01002 AS, TOTAL

01005 BA, DISS

01007 BA, TOTAL

21310 BE, DISS

C1012 BE, TOTAL

01025 CD, DISS

C1027 CD. TOTAL

01030 CR, DISS

01032 CR HEX

01034 CR. TOTAL

01035 CO, DISS

01037 CO. TOTAL

C1G40 CU, DISS

01042 CU, TOTAL

01045 FE, TOTAL

01046 FE, DISS

01049 PB. DISS

01051 PB. TOTAL

PAGE

5

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ATOMIC ABSORPTION

- 21055 MN. TOTAL
- 01055 MN, DISS
- 01057 TL, DISS
- 01059 TL. TOTAL
- 01060 MO, DISS
- 01052 MO, TOTAL
- 01065 NI, DISS
- 01067 NI, TOTAL
- 01075 AG. DISS
- C1077 AG, TOTAL
- 01085 V. DISS
- C1087 V. TOTAL
- 01090 ZN, DISS
- 01092 ZN, TOTAL
- 01095 SB, DISS
- C1C97 SB, TOTAL
- 01100 SN. DISS
- 01102 SN, TOTAL
- C1105 AL, TOTAL
- 01106 AL, DISS
- 01145 SE, DISS
- 01147 SE, TOTAL

OR:GINAL (Red)

ATOMIC ABSORPTION

Clisc TI. DISS

Cliba TI, TOTAL

01220 CP HEX. DISS

71390 HG. DISS

71900 HG, TOTAL

345 CHROMATEGRAPHY

09032 PENTACHLOROPHENOL

99007 PESTICIDES

39330 ALDRIN 39330 DIELDRIN 39360 DDD

39360 000 39365 205 3937000T

39410 HEPTACHLOR 39350 CHLORDANE

THIS LIST MUST BE CONSPICUOUSLY DISPLAYED WITH THE PERMANENT CERTIFICATE AT THE LABORATORY

 W_{a}

AR30400907/33

DEPARTMENT OF ENVIRONMENTAL RESOURCES

OFFICE OF ENVIRONMENTAL PROTECTION

BUREAU OF LABORATORIES

COMMONWEALTH OF PENNSYLVANIA

Certifies that

E. T. C. Corp. 284 Raritan Center Parkway Edison, NJ 08818-7808 I. D. # 68-323

having duly met the requirements of Chapter 109, Subchapter H, Safe Drinking Water Rules and Regulations issued under the Pennsylvania Safe Drinking Water Act of May 1, 1984 (P.L. 206, No. 43), (35 P.S. SS 721. 1-721.17) is hereby appoved as a

Certified Drinking Water Laboratory

To perform the following analyses:

Inorganic Trace Metals, Nitrate/Fluoride

Organic Herbicides/Pesticides, Volatile Organic Chemicals (Group 1 & 2)

Ci transferents Si on revocation To Be Conspicuously Disolared at the Laboratory

AR304

Expiration Date: 7/1/89

The state of the s

Mark M. McClellan

Deputy Secretary for Environmental Protection

H. S. Carlotte

E87074

NICKEL

LEAD

Env Test: 3 & Cert. Corp.

P.O. Box :08

Edison NJ 08818-7808

IATION REPORT

DATE: 11/17

! NUMBER WP019

| ANALYTES | SAMPLE Number | REPORT VALUE | TRUE VALUE* | ACCEPTANCE LIMITS | VARNING LIMITS | PERFORMANCE EVALUATION |
|-----------|------------------|-----------------|----------------|----------------------|-------------------|---------------------------|
| | ETALS IN MIC | | | | | |
| ALUNINUN | 1 | 89.1 | 78.0 | 49.5- 148. | 62.0- 136. | ACCEPTABLE |
| | Z | 877 | 8>5 | 658TUOU. | 707 997. | ACCEPTABLE |
| ARSENIC | 1 | 29.1 | 26.0 | 17.3- 34.1 | 19-4- 32.0 | ACCEPTABLE |
| | 2 | 141 | 130 | 95.3- 161. | 104 153. | ACCEPTABLE ACCEPTABLE |
| BERYLLIUM | 1 | 87.5 | 89.9 | 75-7- 103. | 79-2- 99.6 | ACCEPTABLE |
| | Š | 266 | 270 | 231 306. | 241 296. | ACCEPTABLE |
| CADMIUM | 1 | 19.2 | 10.0 | 7.22- 12.3 | 7.92- 12.1 | ACCEPTABLE |
| | 2 | 152 | 150 | 128 170. | 133 165. | ACCEPTABLE ACCEPTABLE |
| COBALT | 1 | 51.0 | 47.5 | 37.0- 57.4 | 39.6- 54.8 | ACCEPTABLE |
| | 2 | 614 | 594 | 506 694. | 530 670. | ACCEPT |
| CHROMIUM | 1 | 15.4 | 15.0 | 8.74- 20.2 | 10.2- 18.8 | ACCEPTABLE |
| | 2 | 248 | 240 | 181 287. | 194 274. | ACCEPTABLE |
| COPPER | 1 | 41.0 | 40.0 | 31.6- 47.6 | 33.6- 45.6 | ACCEPTABLE |
| | 5 | 178 | 176 | 152 195. | 157 190. | ACCEPTABL. |
| IRON | | | | | | ACCEPTABLE |
| | 2 | 478 | 420 | 357 471. | 371 457. | NOT ACCEPTABLE |
| MERCURY | 1 | 2.51 | 2.40 | 1.52- 3.21 | 1.73- 3.00 | ACCEPTABLE |
| | 2 | 15.4 | 15.6 | 11.6- 20.1 | 12.7- 19.0 | ACCEPTABL |
| MANGANESE | 1 | 39.0 | 37.8 | 27.8- 46.1 | 30.1- 43.8 | ACCEPTABL |
| | 2 | 152 | 147 | 127 164. | 132 159. | ACCEPTABL |

66.5

294

51.5 179 63.0 46.9- 78.8

280 237.- 322.

50.4 37.2- 64.4

168 140.- 197.

50.9- 74.8

248.- 311.

40.6- 61.0

147.- 190.

ACCEPTABL

ACCEPTABL

ACCEPTABL

ACCEPTABL

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY.
AR304011

DATE: 11/17/87

WATER POLLUTION STUDY NUMBER WP019

| | | | | ACCEPTANCE | WARNING | PERFORMANCE |
|------------|---------------|--------------|--------------|--------------------------|--------------------------|--------------------------|
| IALYTES | NUMBER | | | | LIMITS | EVALUATION |
| TRACE M | ETALS IN MICE | ROGRAMS | PER LIT | ER: | | • = = |
| ELENIUM | 1 2 | 20.7 120 | | | 14.0- 24.1 92.4- 141. | |
| ANADIUM | 1 2 | 64.3 645 | | | 50.5- 74.0 547 693. | ACCEPTABLE ACCEPTABLE |
| INC | 1 2 | 30.6 116 | 30.4 114 | | | ACCEPTABLE ACCEPTABLE |
| YNOHITE | 3 4 | 13.2 37.8 | | | 8.22- 20.4 25.9- 50.4 | ACCEPTABLE ACCEPTABLE |
| ER | 3 4 | 17.9 3.60 | 17.5 3.43 | 13.4" 21.5 2.13" 4.95 | 14.4- 20.4 2.49- 4.60 | ACCEPTABLE ACCEPTABLE |
| HALLIUM | 3 4 | 3.00 27.9 | | | 2.01- 4.39 24.1- 40.2 | ACCEPTABLE ACCEPTABLE |
| OLYBDENUM | 3 4 | 4.30 38.0 | | | 1.52- 7.68 23.2- 45.4 | |
| TRONTIUM | 3 4 | 85.0 18.0 | 91.5 18.3 | 73.7- 107. 14.3- 22.2 | 15.4- 21.1 | ACCEPTABLE ACCEPTABLE |
| ITANIUR | 3 4 | 39.0 156 | | | | ACCEPTABLE ACCEPTABLE |
| MINERAL | S IN MILLIGR | AMS PER | LITER: | CEXCEPT AS N | OTED) | |
| H-UNITS | 3 4 | | | | 3.95- 4.07 8.93- 9.33 | ACCEPTABLE ACCEPTABLE |
| PEC. COND. | 1 | 660 | 659 | 592 732. | 610 714. | ACCEPTABLE |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY.

UMHOS/CM AT 25 C) 2 274 272 245.- 302. 252.- 295. ACCEPTABLE

OFIGINAL (Red)

PERFORMANCE EVALUATION REPORT

DATE: 11/17/6

WATER POLLUTION STUDY NUMBER UP019

| BORATORY: N | l | J | 1 | 13 | 6 | , |
|-------------|---|---|---|----|---|---|
|-------------|---|---|---|----|---|---|

| | | = | | | |
|------------------|------------|--|---|---|--|
| SAMPLE NUMBER | REPORT | TRUE VALUE* | ACCEPTANCE LIMITS | E WARNING LINITS | PERFORMANCE EVALUATION |
| MILLIGR | AMS PER | LITER: | CEXCEPT AS | NOTED) | , |
| | | | | | |
| 1 2 | 409 146 | 39 9 158 | 325 482. 95.9- 217 | . 344 462. . 111 202. | ACCEPTABLE ACCEPTABLE |
| . 1 | 66.2 | 63.0 | 54.7- 74. | 0 57-1- 71.6 | ACCEPTABLE |
| 2 | 1.16 | 0.905 | .700- 1.7 | 8 .835- 1.65 | ACCEPTABLE |
| 1 | 0.515 | 0.520 | .42463 | 5 .451508 | ACCEPTABLE |
| 2 | 18.4 | 17.3 | 14.8- 19. | 8 15.4- 19.2 | ACCEPTABLE |
| 1 | 52.3 | 52.6 | 46.0- 58. | 4 47.5- 56.8 | ACCEPTABLE |
| 2 | 13.4 | 13.7 | 10.8- 16. | 2 11.4- 15.6 | ACCEPTABLE |
| | | | | | |
| _ | | | | | |
| 1 | 60.4 | 55.0 | 49.0- 60. | 4 50.4- 59.0 | CHECK FOR ERROR |
| _ | | , | | | |
| 1 | 123 | 113 | 106 128 | . 103 125. | ACCEPTABLE |
| 2 | 53.3 | 52.7 | 47.1- 57. | 1 48.3- 55.9 | ACCEPTABLE |
| 1 | 2.01 | 2.01 | 1.74- 2.2 | 3 1.80- 2.17 | ACCEPTABLE |
| 2 | 0.22 | 0.247 | .155~ .55 | 7 .178514 | ACCEPTABLE |
| 1 | 76.2 | 74.0 | 60.7- 85. | 5 63.8- 82.4 | ACCEPTABLE |
| 2 | 27.8 | 33.0 | 24.5- 59. | 4 26.3- 57.5 | ACCEPTABLE |
| IN MILLIE | RAMS PET | t LITER: | ; | | |
| 1 | 0.478 | 0.500 | .38361 | 4 .411586 | ACCEPTABLE |
| 2 | 1.97 | 2.00 | 1.59- 2.3 | 8 1.68- 2.28 | ACCEPTABLE |
| | NUMBER | NUMBER VALUE 1 409 2 146 1 66.2 2 1.16 1 0.515 2 18.4 1 52.3 2 13.4 1 20.4 2 11.3 1 60.4 2 10.0 1 123 2 53.3 1 2.01 2 0.22 1 76.2 2 7.8 IN MILLIGRAMS PER | NUMBER VALUE VALUE+ 1 409 399 2 146 158 1 66.2 63.0 2 1.16 0.905 1 0.515 0.520 2 18.4 17.3 1 52.3 52.6 2 13.4 13.7 1 20.4 18.0 2 11.3 10.0 1 60.4 55.0 2 10.0 7.49 1 123 113 2 53.3 52.1 1 2.01 2.01 2 0.22 0.247 1 76.2 74.0 2 27.8 33.0 IN MILLIGRAMS PER LITER: | NUMBER VALUE VALUE* LIMITS HILLIGRAMS PER LITER: (EXCEPT AS 1 409 399 325 482. 2 146 158 95.9- 217. 1 66.2 63.0 54.7- 74.0 2 1.16 0.905 .700- 1.78 1 0.515 0.520 .42463 2 18.4 17.3 14.8- 19.3 1 52.3 52.6 46.0- 58.0 2 13.4 13.7 10.8- 16.3 2 13.4 13.7 10.8- 16.3 1 20.4 18.0 14.9- 21.0 2 11.3 10.0 8.29- 11.0 1 60.4 55.0 49.0- 60. 2 10.0 7.49 4.71- 11.0 1 123 113 106 128 2 53.3 52.1 47.1- 57. 1 2.01 2.01 1.74- 2.2 2 0.22 0.247 .15533 1 76.2 74.0 60.7- 85.2 27.8 33.0 24.5- 39. IN HILLIGRAMS PER LITER: | SAMPLE NUMBER VALUE VALUE ACCEPTANCE LIMITS MILLIGRAMS PER LITER: CEXCEPT AS NOTED) 1 409 399 325 482. 344 462. 2 146 158 95.9- 217. 111 202. 1 66.2 63.0 54.7- 74.0 57.1- 71.6 2 1.16 0.905 .700- 1.78 .835- 1.65 1 0.515 0.520 .424635 .451608 2 18.4 17.3 14.8- 19.8 15.4- 19.2 1 52.3 52.6 46.0- 58.4 47.5- 56.8 2 13.4 13.7 10.8- 16.2 11.4- 15.6 1 20.4 18.0 14.9- 21.0 15.6- 20.2 2 11.3 10.0 8.29- 11.5 8.68- 11.1 1 60.4 55.0 49.0- 60.4 50.4- 59.0 2 10.0 7.49 4.71- 11.6 5.57- 10.8 1 123 113 106 128. 103 125. 2 53.3 52.1 47.1- 57.1 48.3- 55.9 1 2.01 2.01 1.74- 2.23 1.80- 2.17 2 0.22 0.247 .155337 .178314 1 76.2 74.0 60.7- 85.5 63.8- 82.4 2 27.8 33.0 24.5- 39.4 26.3- 37.5 IN MILLIGRAMS PER LITER: 1 0.478 0.500 .383614 .411586 2 1.97 2.00 1.59- 2.38 1.68- 2.28 |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY.

PAGE 3

DATE: 11747/87

ATER POLLUTION STUDY NUMBER UP019

| | SAMPLE Number | | | ACCEPTANCE LIMITS | WARNING LINITS | |
|----------------------|------------------|----------------|-------------|--------------------------|--------------------------|--------------------------|
| | | | | | | |
| DEMANDS IN M | ILLIGRA | MS PER L | ITER: | - ₍₁₎ . | | |
| COD | 1 2 | 134 246 | 150 275 | 118 168. 213 307. | 124 162. 225 295. | ACCEPTABLE ACCEPTABLE |
| TOC. | 1 2 | 57.3 109 | 59.2 109 | 46.8- 74.3 86.8- 128. | 50.4- 70.7 92.2- 122. | ACCEPTABLE ACCEPTABLE |
| PCB'S IN HIC | ROGRAMS | PER LIT | ER: | | | |
| PCB-AROCLOR 1016/124 | 2 1 | 3.11 | 4.57 | 2.01- 6.61 | 2.60- 6.02 | ACCEPTABLE |
| 2CB-AROCLOR 1262 | 2 | 1.89 | 1.86 | 1.18- 2.25 | 1.32- 2.11 | ACCEPTABLE |
| PESTICIDES I | N MICRO | GRAMS PE | R LITER | | | |
| ALDRIN | | 0.693 0.303 | | .225- 1.16 .0833460 | | ACCEPTABLE ACCEPTABLE |
| DIELDRIN | | 0.598 0.209 | | .453- 1.12 .134405 | | |
| DDD | | 0.325 0.820 | | .135565 .419- 1.31 | | |
| DDE | | 0.412 0.135 | | .285920 .0926255 | | |
| DDT | | 0.319 0.709 | | -0879477 -330- 1-07 | | ACCEPTABL ACCEPTABL |
| HEPTACHLOR | | 0.598 | | | .272676 .0824216 | ACCEPTABL ACCEPTABL |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY.

DATE: 11/17/82

DRIGINAL (Red)

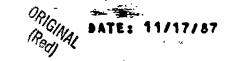
WATER POLLUTION STUDY NUMBER WP019

BORATORY: NJ136

| :ALYTES | SAMPLE Number | REPORT VALUE | TRUE VALUE* | ACCEPTANCE LIMITS | LIMITS | PERFORMANCE |
|---------------------|------------------|-----------------|----------------|--------------------------|--------------------------|---------------------------------|
| PESTICIDES 1 | | | | | * | |
| EPTACHLOR EPOXIDE | 1 2 | 0.086 0.390 | 0.105 0.456 | .0550144 .262603 | .0664432 .305560 | ACCEPTABLE ACCEPTABLE |
| HLORDANE | 3 4 | 6-02 0-629 | 7.73 0.620 | 3.56- 9.39 .240919 | 4.31- 8.65 .327833 | ACCEPTABLE ACCEPTABLE |
| VOLATILE HAI | LOCARBON | IS IN MIC | ROGRAMS | PER LITER; | | |
| ,2 DICHLOROETHANE | 1 2 | 61.9 3.78 | 54.8 3.65 | 37.3- 72.9 .694- 7.74 | 41.9- 68.3 1.60- 6.83 | ACCEPTABLE ACCEPTABLE |
| HLOROFORM | | | | | 62.6- 120. 9.93- 20.0 | |
| ,1,1 TRICHLOROETHA | NE 1 2 | | | 18.4- 52.7 4.84- 15.5 | | ACCEPTA ACCEPTABLE |
| RICHLOROETHENE | 1 2 | 51.4 2.39 | 45.2 2.41 | 30.3- 67.6 1.02- 3.74 | 35.0- 62.8 1.37- 3.39 | ACCEPTABLE ACCEPTABLE |
| ARBONTETRACHLORIDE | 1 2 | | | | | ACCEPTABLE ACCEPTABLE |
| ETRACHLOROETHENE | 1 2 | | | | | S CHECK FOR ERROR Acceptable |
| 3ROMODICHLOROMETHAN | NE 1 | 38.9 8.75 | 32.2 7.24 | 24.5- 45.4 4.11- 11.5 | 27.1- 42.7 5.05- 10.5 | ACCEPTABLE ACCEPTABLE |
| PIBROMOCHLOROMETHAN | NE 1 2 | | 67.7 2.26 | | | ACCEPTABLE ACCEPTABLE |
| BRONGFORM | 1 2 | | | | | 3 ACCEPTABLE 8 ACCEPTABLE |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY.

PAGE 5



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|---|---|---|---|---|---|---|---|---|---|------|---|---|----|
| | | w | ĸ | ~ | • | v | n | | • | - 19 | • | | 30 |

| | AMPLE WHBER | VALUE | VALUE* | LIMI | TS | LIME. | 2.7 | PERFORMANCE EVALUATION |
|----------------------|----------------|---------|---------|---------|------|--------|--------|---------------------------|
| VOLATILE HALO | CARBONS | | | | | | ? ? | |
| METHYLENE CHLORIDE | | | | | | | | ACCEPTABLE |
| | | | | | | | | ACCEPTABLE |
| CHLOROBENZENE | 1 | 31.8 | 30.8 | 18.7- | 43.8 | 21.9- | 40.6 | ACCEPTABLE |
| | 2 | 3.72 | 3.85 | 1.48- | 6.07 | 2.07- | 5.48 | ACCEPTABLE |
| VOLATILE ARON | ATICS I | N MICRO | GRAMS P | ER LITE | R: | | | |
| BENZENE | 4 | 9.58 | 0.80 | 6.20- | 14.0 | 7.29- | 13.n | ACCEPTABLE |
| DENECHE | | | | | | 33.0- | | |
| | | 7 44 | 5 LT | 4 52- | 44 4 | 5 / / | 40.7 | 4665074011 |
| HATBENZENE | 2 | 24.0 | 26.1 | 16.3- | 35.5 | 18.8- | 33.1 | ACCEPTABLE ACCEPTABLE |
| | | | | | | | | |
| TOLUENE | 1 | 5.48 | 5.95 | 3.24- | 8.80 | 3.97- | | ACCEPTABLE |
| | 2 | 27.6 | 29.7 | 20.8- | 39.4 | 23.2- | 37.0 | ACCEPTABLE |
| 1,2-DICHLOROBENZENE | 1 | 5.15 | 5.42 | 1.20- | 9.58 | 2.37- | 8.41 | ACCEPTABLE |
| | | 58.6 | 61.4 | 36.0- | 89.4 | 43.0- | 82.4 | ACCEPTABLE |
| 1,3-DICHLOROBENZENE | . 4 | 3.22 | 3-44 | . 773- | 5_80 | 1-66- | 5-22 | ACCEPTABLE |
| 175 PZCHEORODENZENZ | ż | 24.0 | 26.0 | 10.7- | 38.1 | 14.5- | 34.3 | ACCEPTABL ACCEPTABL |
| | | | | | | | | |
| 1,4-DICHLOROBENZENE | | | 35.8 | | | | | ACCEPT * T |
| | • | 34.0 | 3,00 | | ,,,, | 2310 | 3026 | 7000 |
| MISCELLANEOU | S PARAME | TERS: | | | | | | |
| TOTAL CYANIDE | | | | | | .0805- | | |
| (IN MG/L) | 2 | 0.284 | 0.300 | .174- | .388 | -201- | .361 | ACCEPTABL |
| NON-FILTERABLE RESID | UE 1 | 67-5 | 69.4 | 61-1- | 73.6 | 62.6- | 72.0 | ACCEPTABL |
| (IN MG/L) | Ž | 24.4 | 24.7 | 20.5- | 27.2 | 21.3- | 26.4 | ACCEPTABL |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY. STANDS FOR DETECTION LIMIT

DATE: 11/17/2

WATER POLLUTION STUDY NUMBER WP019

| | ^ | • | \T | Λ | | v | • | M | 14 | 13 | ^ |
|---|---|----|-----------|---|---|---|---|-------|----|----|---|
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(Reg)

| SURATURE: NOTSU | | | | | | |
|---------------------------|------------------|-----------------|----------------|--------------------------|--------------------------|---------------------------|
| ALYTES | SAMPLE Number | REPORT VALUE | TRUE VALUE+ | ACCEPTANCE LIMITS | VARNING LIMITS | PERFORMANCE Evaluation |
| MISCELLAN | EOUS PARAM | ETERS: | | | | |
| IL AND GREASE IN MG/L) | 1 2 | 29.0 10.3 | 35.3 12.8 | 20.9- 43.0 3.99- 18.1 | 23.7- 40.3 5.74- 16.3 | ACCEPTABLE ACCEPTABLE |
| OTAL PHENOLICS [N MG/L) | 1 2 | 0.438 1.16 | | .229775 .588- 1.96 | .298706 .762- 1.79 | ACCEPTABLE ACCEPTABLE |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY.

PAGE 7 (LAST PAGE)

DATE: 07/27/=

WATER SUPPLY STUDY NUMBER WS020

| LABORATORY NJ136 | | | | | |
|------------------|------------------|---------------------|--------------------------|--------------------------|------------------------------|
| ANALYTES | SAMPLE Number | REPORTED VALUE | | ACCEPTANCE LIMITS | PEPFORMANCE EVALUATIONS |
| TRACE META | ALS IN MICPO | GRAMS PER I | LITEP: | | |
| ARSENIC | 1 2 | 109 34.0 | 106 32 _# 0 | 86.8- 121. 25.8- 37.4 | ACCEPTABLE ACCEPTABLE |
| BARIUM | 1 2 | 77•0 746 | | 54.7- 88.6 664 860. | ACCEPTABLE ACCEPTABLE |
| CADMIUM | 1 2 | | | 14.3- 19.6 3.54- 4.79 | ACCEPTABLE NOT ACCEPTABLE |
| CHROMIUM | 1 2 | 13.0 74.5 | 12.7 71.1 | 10.1- 15.6 61.1- 80.9 | ACCEPTABLE ACCEPTABLE |
| LEAD | 1 2 | 26 .1 103 | | 20.6- 30.5 81.7- 113. | ACCEPTABLE ACCEPTABLE |
| ERCURY | 1 2 | 5.14 1.73 | | 3.84- 6.54 1.32- 2.47 | ACCEPTABLE ACCEPTABLE |
| SELENIUM | 1 2 | 9.9 56.3 | 9.71 53.9 | | ACCEPTARLE ACCEPTABLE |
| SILVER | 1 2 | 27.5 15.0 | | 23.1- 31.9 11.2- 16.6 | ACCEPTABLE ACCEPTABLE |
| NITRATE/FI | LUORIDE IN M | ILLIGRAMS | PER LITER | R: | |
| NITRATE AS N | <u>1</u> | 0.948 6.95 | 0.900 7.00 | .762- 1.04 6.18- 7.82 | ACCEPTABLE ACCEPTABLE |
| FLUORIDE | 1 2 | 0.177 1.54 | 0.180 1.60 | .148215 1.48- 1.69 | ACCEPTABLE ACCEPTABLE |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY

PAGE 1



DATE: 07/ //

WATER SUPPLY STUDY NUMBER WS020

| | | | | | PERFORMANCE |
|--------------------------------------|--------|-----------|-----------|------------|----------------|
| ANALYTES | UF3E6 | VALUE | VALUE | LIMITS | 2 PCITAULAV? |
| INSECTICIDES IN | MICRO | GRAMS PER | LITER: | | |
| ENDRIN | 1 | | | | ACCEPTABLE |
| | 2 | | | 3.86- 7.84 | |
| LINDANE | 1 | | ** 0.512 | | |
| | 2 | 4.23 | ** 3-84 | | ACCEPTAPLE |
| METHOXYCHLOR | 1 | 2.37 | 2.22 | 1.34- 3.05 | ACCEPTABLE |
| | 2 | 34.2 | 80.8 | 52.4- 104. | ACCEPTABLE |
| TOXAPHENE | | | | .432- 2.23 | |
| | 4 | 8.93 | 7.09 | 3.85- 9.80 | ACCEPTABLE |
| HERBICIDES IN M | ICROGR | AMS PER L | ITER: | | |
| 2,4-0 | 1 | | | 26-0- 83-8 | |
| | 2 | 3.36 | 3.22 | .413- 5.66 | ACCEPTARL |
| 2,4,5-TP (SILVEX) | | | | 9.42- 41.1 | |
| | 2 | | | 1.23- 5.00 | ACCEPTABLE |
| TRIHALOMETHANES | IN MI | CROGRAMS | PER LITER | t : | |
| CHLOROFORM | 1 | 19.2 | | | ACCEPTABLE |
| | 2 | 54.4 | | | ACCEPTABLE |
| ROMOFORM | 1 | 53.2 | _ | – – | NOT ACCEPTABLE |
| | S | 19.9 | _ | 13.5- 20.3 | ACCEPTABLE |
| BROMODICHLOROMETHAME | 1 | 23.6 | | | ACCEPTABLE |
| · <u>-</u> · · · · · · · · · · · · · | ż | 72.1 | | 50.6- 75.8 | |
| IBROMOCHLOROMETHANE | 1 | 73_2 | 56-9 | 45.5- 68-3 | NOT ACCEPTABLE |
| HINDUC | 2 | 31.2 | | 19.9- 29.9 | |

^{*} BASED UPON THEORETICAL CALCULATIONS, OF A REFERENCE VALUE WHEN NECESSARY SIGNIFICANT GENERAL METHOD BIAS IS ANTICIPATED FOR THIS RESULT.

PAGE 2



DATE: 07/27/

PERFORMANCE EVALUATION REPORT

WATER SUPPLY STUDY NUMBER WS020

| , | | | | |
|---------|-----------|------|--------------|-------------|
| | | | | |
| | | | | |
| SANDIF | REPORTED | TRUE | ACCEPTANCE | PERFORMANCE |
| 381. 20 | KL. OKILD | 1110 | ACCEPTANCE . | FERFORMANCE |

| ANALYTES | NUMBER | VALUE | VALUE+ | | EVALUATIONS |
|-----------------------|----------|----------------|----------------|--------------------------|------------------------------|
| TRIHALOMETHAN | ES IN MI | CROGRAMS P | ER LITER: | | |
| TOTAL TRIHALOMETHANE | 1 2 | 169.2 177.6 | 137.2 154.5 | 110 165. 124 185. | NOT ACCEPTABLE ACCEPTABLE |
| VOLATILE ORGA | NIC COMP | OUNDS IN * | ICROGRAMS | PER LITER: | |
| VINYL CHLORIDE | 1 | 7.36 | 5.98 | 3.59- 3.37 | ACCEPTABLE |
| 1,1-DICHLOROETHYLENE | 1 2 | 3.30 18.3 | 2.53 12.7 | 1.52- 3.54 10.2- 15.2 | ACCEPTABLE NOT ACCEPTABLE |
| 1,2-DICHLOROETHANE | 1 2 | 5.99 11.1 | 6.23 8.90 | 3.74- 8.72 5.34- 12.5 | ACCEPTABLE ACCEPTABLE |
| 1,1,1-TRICHLOROETHANE | 1 2 | 12.6 196 | | 8.40- 12.6 146 219. | ACCEPTABLE ACCEPTABLE |
| ARBON TETRACHLORIDE | 1 | 1.52 | 1.36 | .816- 1.90 | ACCEPTABLE |
| / TRICHLOROETHYLENE | 1 2 | 8.44 10.8 | 8.22 10.3 | 4.93- 11.5 8.24- 12.4 | ACCEPTABLE ACCEPTABLE |
| BENZENE | 1 | 3.76 | 4.32 | 2.59- 6.05 | ACCEPTABLE |
| TETRACHLOROETHYLENE | 2 | 7.60 | 8.16 | 4.90- 11.4 | ACCEPTABLE |
| 1,4-DICHLOROBENZENE | 1 | 7.72 | 6.93 | 4-16- 9-70 | ACCEPTABLE |
| CHLOROBENZENE | 4 | 14.6 | 14.6 | 11.7- 17.5 | ACCEPTABLE |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY

PAGE 3

ORIGINAL (Red)

PERFORMANCE EVALUATION REPORT WATER SUPPLY STUDY NUMBER WS020

DATE: 07/27/

LABOPATORY NJ136

| | | REPORTED VALUE | | ACCEPTANCE LIMITS | PERFORMANC: EVALUATIONS |
|---|----------|-------------------|-----------|----------------------|----------------------------|
| VOLATILE ORGAN | IIC COMP | OUNDS IN M. | ICROGRAMS | PER LITER: | |
| METHYLENE CHLORIDE | 2 | 14-4 | 12.0 | 9-60- 14-4 | ACCEPTABLE |
| 1,1-DICHLOROETHANE | 2 | 11-4 | 10.3 | 8.24- 12.4 | ACCEPTAPLE |
| 1,1-DICHLOROPROPENE | 2 | | 31.6 | 25.3- 37.9 | NOT ACCEPTABLE |
| 1,1,2-TRICHLOROETHANE | 2 | 14.2 | 12.8 | 10-2- 15-4 | ACCEPTARLE |
| 1,1,1,2TETRACHLOROETHA | INE2 | 15.4 | 17.3 | 13.8- 20.8 | ACCEPTABLE |
| 2-CHLOROTOLUENE | 2 | 3.02 | 8.28 | 4.97- 11.6 | NOT ACCEPTABLE |
| -CHLOROTOLUENE | 2 | 3.02 | | D.L D.L. | NOT ACCEPTABLE |
| MISCELLANEOUS | ANALYTE | `S: | | | |
| FURSIDITY (NTU [®] S) | 1 2 | | | | ACCEPTABLE ACCEPTABLE |
| PH-UNITS | 1 | 8.56 | 9.12 | 8.79- 9.34 | NOT ACCEPTABLE |
| SODIUM Milligrams per l iter) | 1 | 13650 | 14.5 | 13.4- 15.9 | NOT ACCEPTABLE |

BASED UPON THEORETICAL CALCULATIONS, OR A REFERENCE VALUE WHEN NECESSARY

PAGE 4 (LAST PAGE)

SIGNIFICANT GENERAL METHOD BIAS IS ANTICIPATED FOR THIS RESULT.

STANDS FOR DETECTION LIMIT

LABORATORY PERFORMANCE AUDITS

EVALUATIONS INCLUDED FOR:

- EVIDENCE AUDIT BY TECHLAW (CLP HEADQUARTERS)
- ON-SITE EVALUATION BY REGION II & LEMSCO (CLP HEADQUARTERS)
- ON-SITE EVALUATION BY NJDEP (X-195 CONTRACT)
- ON-SITE EVALUATION BY NJDEP (CERTIFICATION)

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LABORATORY EVIDENCE AUDIT REPORT

ENVIRONMENTAL TESTING AND CERTIFICATION CORPORATION

March 22, 1988

Environmental Testing and Certification Corporation 284 Raritan Center Parkway Edison, NJ 08818-7808 (201) 225-6792

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¹present at pre-audit meeting 2contacted during audit ³present at post-audit meeting

AR304023 This work was conducted on behalf of the Environmental Protection Agency's (EPA) National Enforcement Investigations Center (NEIC) under EPA Contract #68-01-7369.

INTRODUCTION

An audit of laboratory operations pertaining to laboratory security, sample chain-of-custody, and document control procedures for EPA Dioxin Contract 68-01-7366 (IFB WA 86-K357) was conducted at Environmental Testing and Certification (ETC) Corporation in Edison, New Jersey on March 22, 1988. The audit was conducted by NEIC's Contract Evidence Audit Team (CEAT-TechLaw). Procedures and documentation related to sample receiving, sample storage, sample security, sample tracking, and case file organization and assembly were reviewed for conformance to Evidence Audit Requirements. The results of this audit are discussed in this evidence audit report.

EXECUTIVE SUMMARY

This was the seventh audit of ETC conducted by USEPA representatives in support of the Contract Laboratory Program (CLP). The previous audit was conducted on March 20, 1987 and resulted in no recommendations from the CEAT.

The following five findings (non-conformances to Evidence Audit Requirements) were identified during the present audit and are discussed in this report.

Findings

- 1. The Sample Receipt Form did not contain the name of the laboratory.
- 2. The presence or absence of airbills was not recorded on the Sample Receipt Form.
- 3. The laboratory has not developed written SOPs for sample identification.
- 4. The laboratory has not developed written SOPs for sample tracking.
- 5. Written SOPs for case file preparation did not describe actual procedures used by the laboratory.

As a result of these findings, the following recommendations were made during the debriefing with the laboratory personnel at the conclusion of the audit on March 22, 1988:

Recommendations

- 1. The Sample Receipt Form should be revised to include the name of the laboratory.
- 2. The presence or absence of airbills should be recorded on the Sample Receipt Form.
- 3. The laboratory should develop written SOPs for sample identification.
- 4. The laboratory should develop written SOPs for sample tracking.
- 5. Written SOPs for case file preparation should be revised to describe actual procedures used by the laboratory.

Routine evidence audits will be conducted during the contract period of performance. Corrective action on the above items will be reviewed during the next on-site audit. Periodic audits will be conducted to review continued conformance to Evidence Audit Requirements.

The audit was concluded on March 22, 1988. The audit participants are listed on the cover page of this report.

PROCEDURAL AUDIT

The procedural audit consisted of review and examination of actual and written standard operating procedures (SOPs) and accompanying documents for the following laboratory operations: sample receiving, sample storage, sample tracking (from receipt to completion of analysis), and case file organization and assembly.

Sample Receiving

EPA sample shipments are delivered to the loading dock (Monday - Saturday, 8:00 a.m. - 9:00 p.m.). The designated sample custodian, Bill Deckelmann, signs the airbill and transfers the container to the isolation laboratory. The sample custodian opens the container, inspects the samples, and reviews the shipping documents. Sample receiving information is recorded on the Sample Receipt Form.

During review of the Sample Receipt Form, the auditors observed that the Sample Receipt Form did not contain the name of the laboratory and the presence or absence of airbills was not recorded.

Written SOPs for sample receiving have been developed and implemented and are documented in ETC IFB 2.3.7.8-TCDD SOP Sample Receipt. The auditors read these SOPs, and they accurately describe the procedures in use for sample receiving.

Sample Storage

Dioxin samples and extracts are stored in the locked isolation laboratory. In addition, sample extracts are stored in a small refrigerator located in the gas chromatograph/mass spectrometer (GC/MS) laboratory. Samples are identified with job numbers (unique to sample) and log link numbers (identifies a sample shipment). Extracts are identified with the job number, the type of analysis, and the date. The laboratory maintains the identity of the sample during preparation by writing the sample identifier on the glassware.

Laboratory security is maintained by keeping all access doors locked. Visitors must sign a logbook in the reception area, receive a visitor's badge, and are escorted through the laboratory. Laboratory personnel run a magnetic card through a reader on the receptionist's desk for laboratory entry.

Written SOPs for sample storage and security have been developed and implemented and are documented in <u>ETC IFB 2,3,7,8-TCDD SOP Sample Storage</u>. The auditors read these SOPs, and they accurately describe the procedures in use for sample storage and security. Written SOPs for sample identification have not been developed.

Sample Tracking

Samples may be tracked through the laboratory from receipt to completion of analysis by using the following documents:

- 1. Sample Receipt Form
- 2. Sample Log-In Form
- 3. Laboratory Chronicle: TCDD Extraction
- 4. Laboratory Chronicle: GC/MS Department

The Sample Receipt Form and the Sample Log-In Form are used to record sample receiving information. The Laboratory Chronicles are used to record preparation and analysis information.

Written SOPs for sample tracking have not been developed.

Case File Organization and Assembly

Case files are stored in the document control room. Case files are arranged by EPA case number. Tracy Fedosh or Lori Handle are responsible for case file organization. According to June Baker, QA coordinator, the laboratory has not received confidential documents.

Written SOPs for case file organization and assembly have been developed and implemented and are documented in <u>ETC IFB 2.3.7.8-TCDD SOP Integration of PCDX/PCDF</u>. The auditors read these SOPs, and they do not describe how case file documents are numbered, inventoried, and purged.

EVIDENCE AUDIT

The evidence audit consisted of review and examination of case file documentation. Case files contain the following types of documents:

- 1. Document Inventory
- 2. Airbill
- 3. CLP Dioxin Shipment Record
- 4. Chain-of-Custody Receipt of Cooler
- 5. Chain-of-Custody Record

(Red)

7. ETC Lead Report Tracking Form

8. GC/MS Data - Narrative

9. Final Report - Data, Logs, Etc.

The case file examined during the audit was #8600.

Documentation in the case file is organized and developed according to Evidence Audit Requirements.

AUDIT FINDINGS

The following five findings (non-conformances to Evidence Audit Requirements) are based on the results of the procedural and evidence audits.

- 1. The Sample Receipt Form did not contain the name of the laboratory.
- 2. The presence or absence of airbills was not recorded on the Sample Receipt Form.
- 3. The laboratory has not developed written SOPs for sample identification.
- 4. The laboratory has not developed written SOPs for sample tracking.
- 5. Written SOPs for case file preparation did not describe actual procedures used by the laboratory.

SUMMARY

At the conclusion of the audit on March 22, 1988, a debriefing was held by the audit team with ETC personnel. During this debriefing, the evidence auditors made the following recommendations based on the findings discussed in this report.

- 1. The Sample Receipt Form should be revised to include the name of the laboratory.
- 2. The presence or absence of airbills should be recorded on the Sample Receipt Form.
- 3. The laboratory should develop written SOPs for sample identification.
- 4. The laboratory should develop written SOPs for sample tracking.

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5. Written SOPs for case file preparation should be revised to describe actual procedures used by the laboratory.

April 18, 1988

Mr. Angelo Carasea
Project Officer (WH-548A)
USEPA Headquarters
Office of Solid Waste and
Emergency Response
Analytical Operations Branch
401 M Street, S.W.
Washington, DC 20460

RE: Transmittal of CEAT Laboratory Evidence Audit Report for Environmental Testing and Certification Corporation

Dear Angelo:

Enclosed is a copy of the Contract Evidence Audit Team (CEAT-TechLaw) evidence audit report for the audit conducted at Environmental Testing and Certification (ETC) Corporation on March 22, 1988.

Based on the results of the audit and examination of the audit documentation and procedures used, the chain-of-custody, document control, and evidence security procedures followed by ETC meet or exceed Evidence Audit Requirements. Exceptions to this statement are expressed as findings in the attached report.

CEAT-TechLaw has conducted a management review of the audit report and audit workpapers. The review was made in accordance with generally accepted evidence auditing standards and included such tests of the documentation and other such auditing procedures as were considered necessary in the circumstances.

The subject evidence audit report has been received and approved by NEIC, and copies have been transmitted to the Regional Deputy Project Officer and to the laboratory.

Mr. Angelo Carasea Page Two April 18, 1988

If you have any questions, please contact the Project Officer, Rob Laidlaw, or Don Roche at (303) 236-5122, FTS 776-5122.

Sincerely,

Jeffrey C. Worthington

Contract Evidence Audit Team

Concurrence:

Donald J. Roche

National Enforcement Investigations Center

lkl

Enclosure

cc: Lou Bevilacqua, USEPA Region II DPO

IF: 111-001



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

OFFICE OF RESEARCH AND DEVELOPMENT ENVIRONMENTAL MONITORING SYSTEMS LABORATORY-LAS VEGAS P.O. BOX 93478 LAS VEGAS, NEVADA B9193-3478 (702/798-2100 - FTS 545-2100)

. Gred)

APR 1 5 1988

SUBJECT:

On-Site Laboratory Evaluation Report

FROM:

Jimmie D. Petty

Chief, Quality Assurance Research Branch,

TO:

Angelo Carasea

Organic Project Officer, OERR (WH-548A)

Attached is the routine organic analysis on-site laboratory evaluation report for Environmental Testing Certification (ETC), Edison, New Jersey. The evaluation was conducted on March 22, 1988.

Please contact me at FTS 545-2381 if additional information is needed.

Attachment

~~•

Louis Bevilacqua, Region 2
Jack Farrell, ETC

Environmental Programs Office 1050 E. Flamingo Road, Suite 120, Las Vegas, Nevada 89119 (702) 734-3200

March 30, 1988

United States Environmental Protection Agency P.O. Box 93478 Las Vegas, Nevada 89193-3478

ATTENTION:

DR. J. D. PETTY

VIA:

D. C. PUDVAH & C Kushish \$/6/88

SUBJECT:

ORGANIC ON-SITE LABORATORY EVALUATION REPORT

Dear Dr. Petty:

An Organic On-Site Laboratory Evaluation of Environmental Testing Certification (ETC) performed on March 22, 1988, has been completed. Presently, ETC does not hold an organic contract. The facilities and laboratory procedures were reviewed and suggestion were made in the event of a contract being awarded. The following items must be given attention in order to improve data integrity:

- 1. Volatile and semi-volatile samples and extracts were not kept separate while in cold storage.
- 2. Solvent levels on vials of spiking and calibration solutions should be marked to note any loss resulting from storage.
- 3. All logbooks should be reviewed, signed and dated by supervisory personnel.
- 4. All primary standards must be traceable to EPA reference standards.

DR. J. D. PETTY ORGANIC ON-SITE LABORATORY EVALUATION REPORT PAGE II

ARIGINAL.

Details of the above items may be found in the summary text of this report. An evidentiary audit was conducted simultaneously by the Contract Evidence Audit Team (CEAT) Techlaw.

Very truly yours,

Lisa J. Contreras

Associate Scientist Methods Performance Monitoring Section

LJC/ahh

ec: QA - 3-183

J.O. 70.02 WP-2266C

DAIGINAL BOND

| Laboratory: Environmenta | l Testing and Certification (ETC) | | | | | |
|--------------------------|--|--|--|--|--|--|
| Address: 284 Raritan | Center Parkway | | | | | |
| City: <u>Edison</u> S | tate: NJ Zip: 05818-7308 Telephone: (201) 225-5600 | | | | | |
| | +++++++++++++++++++++++++++++++++++++ | | | | | |
| Type of Evaluation: Orga | nic On-Site Evaluation | | | | | |
| Date of Evaluation: Marc | h 22, 1987 | | | | | |
| Contract Number: Not | Applicable | | | | | |
| | +++++++++++++++++++++++++++++++++++++ | | | | | |
| <u>P</u> I | ERSONNEL CONTACTED | | | | | |
| Name | <u>Title</u> | | | | | |
| Jack Farrell | Technical Manager | | | | | |
| June Baker | QA Coordinator | | | | | |
| Jim Ploscyca | QA Auditor | | | | | |
| Leslie Clarke | Project Representative | | | | | |
| Ken Hebel | Operations Manager | | | | | |
| LABOR | ATORY EVALUATION TEAM | | | | | |
| Name | <u>Title</u> | | | | | |
| Lisa Gatton-Vidulich | Acting DPO, Region 2 | | | | | |
| Stelios Gerazeunis | DPO Representative, Region 2 | | | | | |
| Richard Flotard | Principal Scientist, LEMSCO | | | | | |
| Lisa Contreras | QA Evaluator, LEMSCO | | | | | |
| James Short | Evidence Auditor, Techlaw | | | | | |
| Teri Goldbarg | Ruidence Auditor Techlaw | | | | | |

Summary of Laboratory Evaluation

A. Procedural Changes the Laboratory Should Implement

The following comments refer to deficiencies noted in the Laboratory Evaluation Checklist (Attachment 1).

CONTRACTUAL ITEMS

- Resumes must be submitted to document the qualifications of laboratory personnel.
- 2. Primary standards must be traceable to EPA reference standards. The laboratory must create an SOP for traceability of standards.
- 3. VOA holding blanks should be utilized to determine contamination.

NONCONTRACTUAL ITEMS

- Volatile and semi-volatile samples and extracts should be separated while in cold storage.
- 2. All logbooks should be reviewed, signed, and dated by supervisory personnel.
- 3. Solvent levels on vials of spiking and calibration solutions should be marked to note any loss resulting from storage.
- 4. The SOP for receipt and storage should document actions taken in a problem situation.
- 5. The air-flow of the hoods should be checked and recorded each quarter.
- 6. The balances should be calibrated in the approximate range of sample weight.
- 7. All analytical reagents should be dated upon receipt to assure first-in first-out use.
- 8. The laboratory should use proper correction methods in logbooks.
- 9. The laboratory should create quality control charts available for on-site laboratory inspection.

Attachment 1

Laboratory Evaluation Checklist

I. Organization and Personnel (Page 1 of 2)

| ITEM | YES | NO | COMMENT |
|---|-------------------|----|--------------------|
| Laboratory or Project Manager (individual responsible for overall technical effort) Name: Jack Farrell | x | | Qualified. |
| GC/MS Operator: Name: Tom Rusowich Name: Sam Gibson (Exhibit A, page 8, item E, 10/86) | x | | Qualified. |
| GC/MS Spectral Interpretation Specialist Name: Tom Rusowich/Sam Gibson (Exhibit A, page 8, item E, 10/86) | x | | Qualified. |
| Purge and Trap Specialist Name: Richard Losche Name: (Exhibit A, page 8, item E, 10/86) | | | Resume to be sent. |
| Pesticide Residue Analysis Specialist Name: John Strain Name: (Exhibit A, page 8, item E, 10/86) | | | Resume to be sent. |
| Extraction Concentration Specialist Name: Karen Albertsen Name: (Exhibit A, page 8, item E, 10/86) | x | | Qualified. |

I. Organization and Personnel (Page 2 of 2)

| ITEM | YES | NO | COMMENT |
|---|------------------|----------------------------|---------------------|
| Is the sample custodian designated? If yes, name of sample custodian Name: Bill Deckelmann | x | | , |
| Is the glassware technician designated? Name: Marge Fenyar Name: Anna Stensler | x | ****** ***** ***** ***** * | |
| Was the Quality Assurance Officer available during the evaluation? Name: _Jim Ploscyca | x | | |
| Does the Laboratory Quality Assurance Officer report to senior management levels? | × | | |
| Do personnel assigned to this project have the appropriate educational background to successfully accomplish the objectives of the program? | x | | See above comments. |
| Is the organization adequately staffed to meet project commitments in a timely manner? | × | | |
| Were all key laboratory personnel available? If not list those not available. | x | | |

| Addi | tio | nal C | ommen | ts | |
|------|-----|-------|-------|----|--|
| | | | | | |

| ITEM | YES | NO | COMMENT |
|--|------------------|---------------------|---|
| Are written Standard Operating Procedures (SOPs) developed for receipt and storage of samples? | x | | See comment 1. |
| Is the appropriate portion of the SOP available to the sample custodian at the sample receipt/ storage area? | x | | |
| Are the sample shipping containers opened in a manner which prevents possible laboratory contamination? | x | | Opened in hoods. |
| Are samples that require preservation stored in such a way as to maintain their preservation? VOA-Exhibit D, Pg VOA D-4, Part A, Section 1.1 SVOA-Exhibit D, Pg SV D-4, Part A, Section 1.1 Pest-Exhibit D, Pg Pest D-5 Part A, Section 1.1 | x | | |
| Are volatile samples stored separately from semi-volatile samples? | | x | In sample receipt area. |
| Are VOA holding blanks utilized at a frequency consistent with IFB requirements and is the data maintained for on-site inspection? (VOA-Exhibit D, Pg VOA D-14, Section 2.2) | | x | Had holding blanks when contract in progress. |
| Are adequate facilities provided for storage of samples, including cold storage? | × | | |
| Is the temperature of the cold storage recorded daily in a logbook? | x | | |
| Are temperature excursions noted and are appropriate actions taken when required? | x | | |

II. Sample Receipt and Storage Area (Page 2 of 2)

| ITEM | YES | NO | COMMENT |
|---|--------------|----|----------------|
| Are the sample receipt/storage and temperature logbooks maintained in a manner consistent with GLP? | x | | |
| Has the supervisor of the individual maintaining the document(s) personally examined and reviewed the document(s) periodically, and signed his/ her name therein, together with the date and appropriate comments as to whether or not document(s) are being maintained in an appropriate manner? | - | x | See comment 2. |

Additional Comments

EOP

- 1. The SOP for receipt and storage does not document procedures for a problem situation.
- 2. Logbooks are not reviewed, signed and dated by the supervisor.

. ...

III. Sample Preparation Area (Page 1 of 5)

When touring the facilities, give special attention to: (a) the overall appearance of organization and neatness, (b) the proper maintenance of facilities and instrumentation, (c) the general adequacy of the facilities to accomplish the required work.

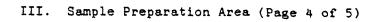
| ITEM | YES | NO | COMMENT |
|--|-------------------|------|------------------------------|
| Is the laboratory maintained in a clean and organized manner? | x | | Except balance. |
| Does the laboratory appear to have adequate workspace (120 sq. feet, 6 linear feet of unencumbered bench space per analyst)? | x | | |
| Are the toxic chemical handling areas either a stainless steel bench or an impervious material covered with absorbent materials? | x | | |
| Are contamination-free areas provided for trace level analytical work? | x | | Adjacent lab. |
| Are contamination-free work areas provided for the handling of toxic materials (e.g., glove box)? | x | | |
| Are exhaust hoods provided to allow contamination-free work with volatile materials? | x | | |
| Is the air flow of the hoods periodically checked and recorded (i.e., once per quarter)? | | × | Not documented this quarter. |
| Are chemical waste disposal policies/procedures well-defined and followed by the laboratory? | x | | |



| ITEM | YES | NO | COMMENT |
|--|--------------|------------------|----------------------|
| Can the laboratory supervisor document that trace-free water is available for preparation of standards and blanks? | | x | See comment 7. |
| Is the analytical balance located away from draft and areas subject to rapid temperature changes? | x | | |
| Has the balance been calibrated and checked within one year by a certified technician? | x | | |
| Are the balance(s) routinely checked with the appropriate range of class S (traceable) weights before each weighing session and are the results recorded in a logbook? | - | x | See comment 3. |
| Are the solvent storage cabinets properly vented as appropriate for the prevention of possible laboratory contamination? | # # # | x | Not vented. |
| Are reagent grade or higher purity chemicals used to prepare standards? | x | | |
| Are analytical reagents dated upon receipt? | | x | Reagents not dated. |
| Are reagent inventories maintained on a first-in, first-out basis? | x | | See comment 4. |
| Are analytical reagents checked out before use? | x | | |

exef)

| ITEM | YES | NO | COMMENT |
|---|---------------------------------------|-------------------|--|
| Are spiking/calibration standards preparation and tracking logbook(s) maintained for: Base-neutral/acids (Exhibit E, Pg 8, Section 8) (Exhibit D, Pg SV D-6, Section 4.7) Pesticides (Exhibit E, Pg 8, Section 8) (Exhibit D, Pg Pest D-8, Section 4.7) Volatiles (Exhibit E, Pg 8, Section 8) (Exhibit E, Pg 8, Section 8) (Exhibit D, Pg VOA D-18, Section 4.6) | x x x x x x x x x x | | Not using acceptable correction methods. |
| Are the primary standards traceable to EPA reference standards for: (Exhibit E, Pg 6, Section 5.1.3) Base-neutral/acids (Exhibit D, Pg SV D-26, Section 3.2) Pesticides (Exhibit D, Pg Pest D-32, Section 4.2.1) Volatiles (Exhibit D, pg VOA D-17, Section 4.4) | | <u>x</u> <u>x</u> | SOP must be written. |
| Are fresh analytical standards prepared at a frequency consistent with the IFB requirements for: Base-neutral/acids (Exhibit D, Pg SV D-27, Section 3.2.1.2) Pesticides (Exhibit D, Pg Pest D-32, Section 4.2.2) Volatiles (Exhibit D, Pg VOA D-18, Section 4.4.5) | | | |
| Are reference materials properly labeled with concentrations, date of preparation, and the identity of the person preparing the sample, and/or is a traceable reference code number used? | × | | |





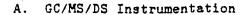
| ITEM | YES | NO | COMMENT |
|---|---------|----|------------------------|
| Do the analysts record bench data in a neat and accurate manner? | × | | |
| Are the sample preparation area and temperature logbooks maintained in a manner consistent with GLP? | x | | |
| Has the supervisor of the individual maintaining the document(s) personally examined and reviewed the document(s) periodically, and signed his/her name therein, together with the date and appropriate comments as to whether or not the document(s) is being maintained in an appropriate manner? | | x | See comment 2. |
| Are standards stored separately from sample extracts? | x | | |
| Are volatile and semi-volatile solutions properly segregated? | | x | Not in sample receipt. |
| Is the appropriate portion of the SOP available to the analyst at the sample preparation area? | x | | |
| Is the SOP for glassware washing posted at the cleaning station? | x | | |
| Is the temperature of the refrigerators/freezers recorded daily? | × | | |
| Are temperature excursions noted and appropriate actions taken when required? | × | | |

III. Sam-le Preparation Area (Page 5 of 5)

Additional Comments

- The balance should be calibrated in the approximate range of sample weight.
- 4. Laboratory management stated reagents were used on first-in, first-out basis.
- 7. The laboratory does not document trace free water.

IV. Sample Analysis Instrumentation (Page 1 of 6)



| | | | | | | Purg | e and | Trap |
|---------------|-----|--------------|-------|-----------------------|----------------------|-----------------|-------|---------|
| | | Manufacturer | Model | Software/ Revision | Installation Date | Manuf. Model | ID # | Install |
| GC/MS ID # | С | нр | 5995 | Rev. E | * | TEK LSC 2 | | * |
| GC/MS ID # | , G | НР | 5995 | Rev. E | ¥ | | | |

^{*} The installation date was unavailable during the on-site. The laboratory will send this information.

| ITEM | YES | NO | COMMENT |
|---|---------|----------|-------------------|
| Are manufacturer's operating manuals readily available to the operator? | x | | |
| Is service maintenance by contract? | x | | |
| Are extensive in-house replacement parts available? | x | | |
| Is the preventative maintenance applied? | x | | |
| Is a permanent service record maintained in a logbook? | × | | |
| Has the instrument been modified in any way? | | x | No modifications. |
| Is the instrument properly vented or are appropriate traps in place? | x | | |
| Is a glass jet separator in place and operational? | x | | |
| Is a split/splitless capillary injector in place? | x | | |
| Is raw data being archived properly (i.e., magnetic tape)? | x | ! | |



IV. Sample Analysis Instrumentation (Page 3 of 6)

| ITEM | YES | NO | COMMENT |
|---|-------------------|---------------------------------------|----------------|
| Are in-house quality control charts maintained and available for on-site inspection? Base-neutral/acids: EICP areas of internal standards Retention times of internal standards (Exhibit E, Pg 41, Section 6.1.1.1) Volatiles: EICP areas of internal standards Retention times of internal standards (Exhibit E, Pg 23, Section 6.1.1.1) | | x x x x x x x x x x | See comment 5. |
| Are the corrective actions described in the IFB implemented and documented as required? Base-neutral/acids: Volatiles: (Exhibit E, Pg 23 and 41, Section 6.1.1.1) | <u>x</u> <u>x</u> | | - |

Additional Comments

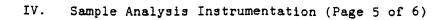
5. The program to create quality control charts is available to the QA officer, but the laboratory is not producing the charts as of yet.

াংস্ক্রা, Sample Analysis Instrumentation (Page 4 of 6)

B. GC Instrumentation

| | | | | Data System | | | |
|--------|-----------------|----------|----------------------|-----------------|----------------------|--|--|
| | Manufacturer De | etectors | Installation Date | Manuf. Model | Installation Date | | |
| GC/ | Hewlett-Packard | 200 | * | HP 1000 | * | | |
| ID # N | 5890 | BC2 | | | | | |
| GC/ | Hewlett-Packard | | | HP | | | |
| ID # B | 5890 | EC2 | * | 1000 | * | | |
| GC/ | Hewlett-Packard | | | НР | | | |
| ID # K | 5880 | EC2 | * | 1000 | * | | |
| GC/ | Hewlett-Packard | | | HP | | | |
| ID # H | 5880 | EC2 | * | 1000 | * | | |

^{*} The installation date was unavailable during the on-site. The laboratory will send this information.





| ITEM | YES | NO | COMMENT |
|--|-------------|-----|---------------------------------|
| Are the manufacturer's operating manuals readily available to the operator? | x | | |
| Is service maintenance by contract? | x | | |
| Are in-house replacement parts available? | × | | |
| Is preventative maintenance applied? | x | | |
| Is a permanent service record maintained in a logbook? | x | | |
| Has the instrument been modified in any way? | | × | No modifications. |
| Is the instrument properly vented or are appropriate traps in place? | x | x | See comment 6. |
| Are Arochlor 1221 and 1232 standards run at the proper frequency and the data maintained for on-site inspection? (Exhibit E, Pg 55, Section 4.3.4.2) | | | Not applicable. |
| Are data generated by the Alumina Equivalency Check available for on-site inspection? If yes, are the following criteria met? (Exhibit D. Pg 15, Section 1.5.8) | | | Not applicable. |
| Is the absence of tribromophenol noted? | | - i | Not applicable. |
| Is the percent recovery of all single componnent pesticides \geq 80%, except for endosulfan sulfate which must be \geq 60%, and endrin Aldehyde which should not be recovered? | | | Not applicable. |

- IV. Sample Analysis Instrumentation (Page 6 of 6)
 - C. Additional Comments
- 6. The purge was not trapped.

V. Data Handling and Review



| ITEM | YES | NO | COMMENT |
|---|-----|-------------------|--|
| Are data calculations spot-checked by a second person? | × | - | l sample/batch or 10 percent of samples. |
| Do records indicate that appropriate corrective action has been taken when analytical results fail to meet QC criteria? | x | | |
| Are computer programs validated before use? | x | - | In-house preparation. |
| Do supervisory personnel review the data and QC results? | x | | |

| ITEM | YES | NO | COMMENT |
|--|--------------|------|---------|
| Does the laboratory maintain a project specific Quality Control Manual? | x | | |
| Are outdated portions of the QC Manual properly archived? | x | | |
| does the manual address the important elements of a QC program, including the following: | | | |
| a. Personnel? | × | | |
| b. Facilities and equipment? | x | | |
| c. Operation of instruments? | x | | |
| d. Documentation of procedures? | x | | |
| e. Preventative maintenance? | x | | |
| f. Reliability of data? | x | | |
| g. Data validation? | x | | |
| h. Feedback and corrective action? | × | | |

VII. Summary



A. Summary Checksheet (Page 1 of 2)

| ITEM | YES | NO | COMMENT |
|---|-------------|---------|---------|
| Do responses to the evaluation indicate that project and supervisory personnel are aware of QA/QC and its application to the project? | x | | |
| Do project and supervisory personnel place positive emphasis on QA/QC? | x | | |
| Have responses with respect to QA/QC aspects of the project been open and direct? | x | | |
| Has a cooperative attitude been displayed by all project and supervisory personnel? | x | | |
| Have any QA/QC deficiencies been discussed before leaving? | x | | |
| Is the overall quality assurance adequate to accomplish the objectives of the project? | x | | |
| Have corrective actions recommended during previous evaluations been implemented? If not, provide details in Section VII.B. | | x | |

B. Additional Comments

- 1. Volatile and semi-volatile samples and extracts were not kept separate while in cold storage.
- 2. Solvent levels on vials of spiking and calibration solution should be marked to note any loss resulting from storage.
- 3. It was recommended that solvents and other reagents be dated upon receipt to assure first-in first-out use.



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

OFFICE OF RESEARCH AND DEVELOPMENT ENVIRONMENTAL MONITORING SYSTEMS LABORATORY-LAS VEGAS P.O. BOX 93478 LAS VEGAS, NEVADA 89193-3478 (702/798-2100 - FTS 545-2100) PICINAL

APR 1 5 1988

SUBJECT:

On-Site Laboratory Evaluation Repor

FROM:

Jimmie D. Petty

Chief, Quality Assurance Research Branch, QAD

TO:

Angelo Carasea

Organic Project Officer, OERR (WH-548A)

Attached is the routine organic dioxin analysis on-site laboratory evaluation report for Environmental Testing and Certification, Edison, New Jersey. The evaluation was conducted on March 22, 1988.

Please contact me at FTS 545-2381 if additional information is needed.

Attachment

cc:

Louis Bevilacqua, Region 2 Jack Farrell, ETC



Environmental Programs Office 1050 E. Flamingo Road, Suite 120, Las Vegas, Nevada 89119 (702) 734-3200

7 April 1988

United States Environmental Protection Agency P.O. Box 93478 Las Vegas, NV 89193-3478

ATTENTION:

Dr. J. D. Petty 2 C. Pudvah 4/8/88

VIA:

SUBJECT:

Routine Dioxin Organic On-Site Laboratory Evaluation Report.

For Environmental Testing and Certification on March 22, 1988.

Dear Dr. Petty:

The routine Dioxin Organic On-Site Evaluation of Environmental Testing and Certification has been completed. The following items must be given attention in order to improve data integrity:

- The SOP for the sample receipt area should be expanded to include corrective actions. #5M-0-009 section 4 has been reused to melode priting
- The analyst preparing standards should mark the initial level of the solution on the container.
- Balances used to weigh samples should be calibrated with a weight in the same range as the size of a typical sample aliquot.
- Analytical reagents should be dated upon receipt and opening. The laboratory should document that they have checked the purity of reagents used in these analyses.
- The laboratory should maintain a file to document water quality by keeping a series of method blanks in a folder available for on-site inspection.
- All logs associated with this project must be periodically reviewed by a supervisor or his designee, signed and dated, along with comments on the acceptability of the document.

Dr. J. D. Petty ROUTINE DIOXIN ORGANIC ON-SITE LABORATORY EVALUATION REPORT FOR ETC. Page $2\,$



Details of the above items may be found in the text of this report. An evidentiary audit was conducted simultaneously by the Contract Evidence Audit Team (CEAT) Techlaw. Their findings will be provided in a separate report.

Very truly yours,

P. E. Flotand
R. D. Flotand

Principal Scientist

Quality Assurance Department

ndf Attachment

cc: QA 4-110 J.O. 70.02 Laboratory: Envisapmental Testing and Certification

Address: 284 Raritan Center Parkway

City: Edison State: N.J. Zip Code: 05818-7308 Telephone: 201-225-5600

Type of Evaluation: Routine Dioxin Organic On-Site Laboratory Evaluation

Date of Evaluation: 22 March 1988

Contract Number: 68-81-7366

Contract Title: Chemical Analytical Services for Dioxin

Personnel Contacted:

<u>Name</u>

Jack Farrel
Dave Speis
June Baker
Ken Hebel
Jim Ploscyca
Leslie Clarke

Title:

Technical Manager GC/MS Manager QA Coordinator Operations Manager QA Auditor Project Representative

<u>Laboratory Evaluation Team:</u>

Name

Richard Flotard Lisa Contreras Lisa Gatton-Vidulich Stelios Gerazeunis James Short Teri Goldberg

Title:

Principal Scientist, LEMSCO Associate Scientist, LEMSCO Acting DPO. USEPA Region II, USEPA Staff Associate, TECHLAW Associate Consultant, TECHLAW



Summary of Laboratory Evaluation

A. Procedural Changes the Laboratory Should Implement



The following comments refer to the deficiencies noted in the Laboratory Evaluation Checklist (Attachment 1)

CONTRACTUAL ITEMS

- 1. The laboratory must submit current resumes for all employees added to this project since the last on-site evaluation.
- The SOP for the sample receipt area should be expanded to include corrective actions.

NONCONTRACTUAL ITEMS

- 1. The analyst preparing standards should mark the initial level of the solution on the container.
- 2. Balances used to weigh samples should be calibrated with a weight in the same range as the size of a typical sample aliquot.
- 3. Analytical reagents should be dated upon receipt and opening. The laboratory should document that they have checked the purity of reagents used in these analyses.
- 4. The laboratory should maintain a file to document water quality by keeping a series of method blanks in a folder available for on-site inspection.
- 5. The laboratory must submit information documenting instrument installation dates to EMSL-LV for GC/MS instruments used for this task.
- 6. All logs associated with this project must be periodically reviewed by a supervisor or his designee, signed and dated, along with comments on the acceptability of the document.
- B. Review of Data Audit Report

The following comments refer to the Summary Conclusion section of the data audit report for SAS Case 8600 (Attachment 2).

Five minor errors were noted in the audit, for an overall score of 0.5 operational defects.

Report
Item No.

Comments

Action*

m-2 Incorrect formula used to calculate SD and has been therefore RSD (used N instead of N-1 in the corrected denominator of the formula)

see audit The laboratory experienced problems with enclosure the column performance solution. E. Kantor case of EMSL-LV said that other labs were not having narrative a problem with this solution.

B. Review of Data Audit Report, Continued

Recort
Item No.

Comments

The laboratory used a six month old initial calibration in this case. Even though contract requirements were met, sensitivity enclosure of the GC/MS had decreased significantly and pages 6, 7 good laboratory practice would indicate the

Action*
has been

corrected

see audit enclosure page 3 Concentration calibration solutions used in this SAS were different from what is listed in the RAS contract. The laboratory did not explain why the standard CC solutions were not used.

need to determine why this is happening.

3

C. Issues to be Resolved by the Project Officer/Deputy Project Officer (PO/DPO):

No additional problems were noted.

- * = 1. No action required
 - 2. Resubmission Required
 - 3. Action Required by Project Officer

Attachment 1

Laboratory Evaluation Checklist



I. Organization and Personnel (page 1 of 2)

| ITEM | YES | 100 | COMMENT |
|--|-------------------|-------|---|
| Laboratory or Project Manager (individual responsible for overall technical effort) | | 1 | |
| Name: Jack Farrell | X | ! | Qualified |
| GC/MS Operator Name: Tom Rusowich Name: Sam Gibson Experience: 1 year minimum requirement per appropriate instrument | × × | • | Qualified Qualified |
| GC/MS Spectral Interpretation Specialist Name: Tom Rusowich Sam Gibson Experience: 2 years minimum requirement | × | ! | - Qualified |
| Extraction Concentration Expert Name: Karen Albertsen Name: Paul Cormier Experience: 6 months minimum requirement | X | : | Qualified Uncertain. resume to be sent |
| Do personnel assigned to this project have the appropriate educational background to successfully accomplish the objectives of the program? | l | 1 | |
| Do personnel assigned to this project have the appropriate level and type of experience to successfully accomplish the objectives of this program? | X | | ! ! ! |
| Is the organization adequately staffed to meet project commitments in a timely manner? | × | | ! ! |

| Τ. | Organization | and | Personnel | (page | 2 | ο÷ | 23 |
|----|---------------|--------|-------------|-------|---|----|-----|
| A | or danization | 2.11 G | i ei sonnei | /hade | ~ | OΤ | ~ / |

ORIGINAL (Red)

| Does the laboratory Quality Assurance Supervisor report to senior management levels? | × | ! ! | . - - | ; ; ; |
|---|--------------|--------|---------------|-------------|
| Was the Project Manager available during the evaluation? | ; × | | · - - | |
| Was the Quality Assurance Supervisor available during the evaluation? | × | | | ; |

Additional Comments: The laboratory was requested to send resumes for all personnel involved with the dioxin and CLP organic programs.

A. General Facilities (page 1 of 2)

Sod)

| | | | 47 |
|--|-------------------|------------------|--|
| I T EM | IYES | 1 011 | COMMENT |
| Does the laboratory appear to have adequate workspace (120 sq. feet, 6 linear feet of unencumbered bench space per analyst)? | × | | |
| Are voltage control devices used on major instrumentation? | × | ! | |
| Does the laboratory have a source of distilled/demineralized water? | | ! ! ! | ; |
| Is the conductivity of distilled/demineralized water routinely checked and recorded? | ; ; ; | | <pre>! Water quality is ! not currently ! being documented ! by ETC. !</pre> |
| Is the analytical balance located away from draft and areas subject to rapid temperature changes? | ; ; × | ! ! | |
| Has the balance been calibrated and checked within one year by a certified technician? | : × | : | ! ! |
| Are the balance(s) routinely checked with the appropriate range of class S (traceable) weights before each weighing session and are results recorded in a logbook? | X | 1 | Balances are checked, but no using weights 1 the actual rang of the samples. |
| Are properly filtered exhaust hoods provided to allow efficient work with hazardous/toxic materials? | × | : : : : | ! ! ! |
| Is the laboratory maintained in a clean and organized manner? | X | | |
| Is a glove box available to allow efficient work with hazardous/toxic materials? | × | | .1 |
| | 1 | 1 | 1 |

A. General Facilities (page 2 of 2)

| I T E M | | | (Red) AL |
|---|-------------|------------|--|
| 1154 | ; 7ES | ! NO ! | COMMENT |
| Is the toxic chemical handling area either a stainless steel bench or an impervious material covered with absorbent material? | | | |
| Are adequate facilities provided for storage of samples, extracts, and calibration standards, including temperature controlled storage? | : : : | | |
| Is the temperature of the cold storage units recorded daily in logbooks? | × | 1 1 | Yes, with the exception of the clean lab unit. |
| Are chemical waste disposal policies/procedures adequate? | × | | |
| Are contamination-free areas provided for trace level analytical work? | X | | |
| Can the laboratory supervisor document that trace free water is available for preparation of standards and blanks? | 1 | ; ; | The laboratory does not document this item. |
| Is the laboratory secure? | X | | |
| Can the laboratory supervisor document that organic solvents used are free of trace contaminants? | | 1 | Documentation was not avail- able to the evaluators. |

Additional Comments on Laboratory Facilities:

- The laboratory has agreed to begin to document the quality of the water used for dioxin analysis by maintaining a file of method blanks for future on-site inspections.
- A temperature log will be prepared for the refrigerator in the dioxin clean laboratory.

B. Equipment (page 1 of 2)

| 1. | GC/MS/DS | Instrumentation |
|----|----------|-----------------|
|----|----------|-----------------|

| | | Manufacturer | Model | Installation Date | Data System |
|---------------|---|-----------------|-----------------------|----------------------|----------------|
| GC/MS ID # | ĸ | Hewlett Packard | . 5390 GC 5920 MSD | NA* | RTE-6 Rev.E |
| GC/MS ID # | 0 | Hewlett Packard | 5996 | NA* | RTE-6 Rev.E |
| GC/MS ID # | J | Hewlett Packard | 5996 | NA* | RTE-o Rev.E |

^{*} Information on the installation of the equipment was not available during the on-site visit. The laboratory has agreed to forward this information to EMSL-LV.

Additional Comments on GC/MS/DS Instrumentation:

None

ORIGINAL (Red)

| ITEM | Y E S | NO ! | COMMENT |
|--|------------------|----------------------------|--------------------------------|
| Are manufacturer's operating manuals readily available to the operator? | × | | |
| Is there a calibration protocol available to the operator? | X | ; ; ; ; ; ; | |
| Are calibration results kept in a permanent record? | × | ! ! ! | |
| Does the laboratory have service contracts for the laboratory instruments? | | i 1 1 1 | |
| Is preventative maintenance applied? | l X | 1 | |
| Is a permanent service record maintained in a logbook? | X | | |
| Has the instrument been modified in any way? | | | No modifications to the units. |
| Is the instrument properly vented or are appropriate traps in place? | × | i : | 1 |
| Is a 9-track magnetic tape unit available? | X | | 1 |
| Is a split/splitless capillary injector in place? | X | : | |
| Is the column direct to the source? | i × | | |
| Are sufficient in-house replacement parts available? | i x | 1 | ! |

Additional Comments on GC/MS Instrumentation:

None

II. Documentation (page 1 of 2)

When reviewing documentation, give special attention to:

- (a) traceability
- (b) neatness and completion

ORIGINAL (Dad)

A. Documentation/Tracking

| ITEM | YES | 1 0/1 | COMMENT |
|---|--|--------|---|
| Is a sample custodian designated? If yes, name of sample custodian. Name: Bill Deckelmann | × | | |
| Are the sample custodian's procedures and responsibilities documented? If yes, where are these documented? | —————————————————————————————————————— | | Documented in the sample receipt SOP. |
| Is a written Standard Operating Procedure (SOP) developed for receipt of samples? If yes, where is the SOP documented (laboratory manual, written instructions, etc.)? | × | 1 | Documented in the QA manual. A copy is kept in the sample receipt area. |
| Are quality assurance procedures documented and available to the analysts? If yes, where are these documented? | | | Located in the QA manual. |
| Are written Standard Operating Procedures (SOP) developed for compiling and maintaining sample document files? If yes, where are the SOPs documented (laboratory manual, written instructions, etc.)? | X | | Located in the QA manual. |
| Are the magnetic tapes stored in a secure area? | × | ! ! | |
| Is a permanently-bound notebook with preprinted, consecutively-numbered pages being used? | | 1 | Computer generated data sheets are used. |

| ITEM | YES | NO : | COMMENT |
|--|---|---------------------------------------|--|
| Is the type of work clearly displayed on the notebook (i.e. EPA Extraction)? | : × | 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 | |
| Is the notebook maintained in a legible manner? | × | : | |
| Are entries noting anomalies routinely recorded? | \ ! | | None was observed. |
| Has the analyst avoided obliterating entries? | : × | | See Note 1. |
| Are inserts (i.e., chromatograms, domputer printout, etc.) permanently affixed in the note-book and signed across insert edge and page? | : | | The use of this technique for enclosures was suggested. |
| Has the supervisor of the individual maintaining the notebook personally examined and reviewed the notebook periodically, and signed his/her name therein, together with the date and appropriate comments as to whether or not the notebook is being maintained in an appropriate manner? | , | : | This practice will be instituted by the laboratory. It is not currently practiced regularly. |
| Where applicable, is the notebook holder referencing reports or memoranda pertinent to the contents of an entry? | : : : | : : | No examples of this were noted in any of the notebooks. |

Note 1. The laboratory should follow prescribed procedures for error correction in laboratory notebooks. Cross out the incorrect entreplace the correction nearby, and sign and date the new entry.

III. Analytical Methodology (page 1 of 2)

| | | | ORIGINAL |
|---|---------------------------------------|---------------------------------------|---|
| ITEM | YES | NO I | |
| Are the required methods used? | × | · · · · · · · · · · · · · · · · · · · | · |
| Is there any unauthorized deviation from contract methodology? | · · · · · · · · · · · · · · · · · · · | ! | None was obser- ved but no samples were being processed. |
| Are written analytical procedures provided to the analyst? | : × | ! ! ! ! | |
| Are distilled-in-glass grade or other high purity chemicals used to prepare standards? | : : × | ! ! | ! ! ! |
| Are fresh analytical standards prepared at a frequency consistent with good QA? | ! ! X | ; | 1 1 1 1 1 |
| Are reference materials properly labeled with concentrations, date of preparation, and the identity of the person preparing the sample? | X | 1 | The initial level of stock standards should be marked on the container. |
| Is a standards preparation and tracking logbook maintained? | X | ! ! | |
| Do the analysts record bench data in a neat and accurate manner? | i : : | 1 | |
| Is the appropriate instrumentation used in accordance with the required protocol? | X | | |



| ITEM | YES | NO I | COMMENT : |
|--|------------------|-----------------------|-------------|
| Does the laboratory maintain a Quality Control Manual? | X | | |
| Does the manual address the important elements of a QC program, including the following? | × | ; ; ; ; ; | |
| a. Personnel? | X | : : : | 1 |
| b. Facilities and equipment? | X | | |
| c. Operation of instruments? | X | ; ; ; | |
| d. Documentation of procedures? | X X | ! ! ! | 1 |
| e. Procurement and inventory practices? | X | ! ! | |
| f. Preventive maintenance? | ; ; × ; | ! ! ! | ! ! |
| g. Reliability of data? | ; X | ! ! ! | |
| h. Data validation? | X | | |
| i. Feedback and corrective action? | X | ! ! ! | |
| j. Instrument calibration? | × | | |
| k. Recordkeeping? | ¦ × | | - - |
| l. Internal audits? | X | ! ! | |

IV. Quality Control Manual Checklist (page 2 of 2)

| ITEM | YES | NO | COMMENT ! |
|--|---------------|--------|------------------|
| Are QC responsibilities and reporting relation- ships clearly defined? | X | 1 | |
| Have standard curves been adequately documented? | 1 | | Not observed |
| Are laboratory standards traceable? |) × | ! ! | 1 |
| Are quality control charts maintained for each routine analysis? | | ! | { } ! |
| Do QC records show corrective action when analytical results fail to meet QC criteria? | | | ! ! |
| Do supervisory personnel review the data and QC results? | ; ; ; × | | |

V. Data Handling Checklist (page 1 of 1)

| ITEM | YES!NO | COMMENT |
|--|--------|---|
| Are data calculations checked by a second person? | • | 10% of the calculations are checked with a minimum of 1 sample per batch. |
| Are data calculations documented? | | 1 |
| De records indicate that corrective action has been taken on rejected data? | | |
| Are limits of detection determined and reported properly? | | |
| Are all data and records retained for the required time? | | * |
| Are quality control data (e.g., standard curve results of duplication and spikes) accessible for all analytical results? | | Not observed by on-site auditor. |

A. Summary Checklist (page 1 of 2)

| ITEM | TYES! | NO 1 | COMMENT |
|---|---------------------------------------|---------------------------------|---|
| Do responses to the evaluation indicate that project and supervisory personnel are aware of QA/QC and its application to the project? | ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; | ; 1 1 1 1 1 1 | |
| Do project and supervisory personnel place positive emphasis on QA/QC? | ; × ; | | |
| Have responses with respect to QA/QC aspects of the project been open and direct? | ; — ; | | |
| Has a cooperative attitude been displayed by all project and supervisory personnel? | ; ; × | | |
| Does the organization place the proper emphasis on quality assurance? | X | ; ; | |
| Have any QA/QC deficiencies been discussed before leaving? | X | 1 1 | |
| Is the overall quality assurance adequate to accomplish the objectives of the project? | ; ; ; × | | |
| Have corrective actions recommended during previous evaluations been implemented? If not, provide details in Section VII.B | - | X | Most have been implemented. Those listed on page 20 have not been done. |
| Are any corrective actions required? If so, list the necessary actions below. | × | 1 | See section A, page 5 for a list of actions |

- B. Summary Comments and Corrective Actions (page 2 of 2)
 - The following items were noted during the on-site and were similar or identical to requests made following the previous on-site evaluation.
 - The analyst preparing standards should mark the initial level of the solution on the container. This same request was noted in the previous on-site evaluation.
- 2. Balances used to weigh samples should be calibrated with a weight in the same range as the size of a typical sample aliquot. In the previous on-site, the use of balance logs had just begun. The laboratory implemented a procedure for checking the calibration of the balanced, but did not implement it correctly.

CRICHTAL ROLL

DIOXIN DATA AUDIT REPORT

| | 8600/DB0355, | | | |
|--|---|--|--|--|
| Laboratory: ETC Corporation | Case/Batch: DB0356 | | | |
| Region: 2 Number of Samples: 22 | IFB/SAS: 68-01-7366 | | | |
| Date Received: 12/17/87 Date | Audited: 2/23-25/88 | | | |
| Contract Revision Date: 9/86 Date R | eviewed: 2/29/88 | | | |
| * Audit Plate revised 6/87. | | | | |
| I. Data Summary Form (B-1) II. Initial Calibration Summary (B-2) III. Routine Calibration Summary (B-3) IV. Quality Control Summary (B-4) V. Other Deliverables VI. Calibration Quality Assurance Criteria VII. Identification Criteria VIII. Native TCDD Spike Results IX. Laboratory Duplicate Analysis Results X. Blank Analysis Results XI. PE Sample Results Total Number of Defects | C=Critical M=Major m=mino 1 | | | |
| This translates into 0.5 Operational Defects Operational Defects = (1.0 x Critical) = (0.3 x Hajor) = (0.1 x minor) | | | | |
| Reviewed by: | Initial Audit by: | | | |
| G. L. Robertson Scientific Supervisor Lockheed Engineering and Management Services Company P.O. Box 15027 Las Vegas, NV 89114 Phone: (702) 734-3326 | Rein forderen. L. J. Contreras Associate Scientist | | | |

SUMMARY COMMENTS/CONCLUSIONS



Soil Samples

| EPA Number | Lab ID |
|------------|--------|
| | |
| MB | |
| DB035514 | |
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| DB035518 | |
| DB035519 | |
| DB035520 | |
| DB035521 | |
| DB035522 | |
| DB035523 | |
| DB035523-N | |
| DB035524 | |
| DB035614 | |
| DB035614-D | |
| DB035615 | |
| DB035616 | |
| DB035617 | |
| DB035618 | |
| DB035619 | |
| DB035620 | |
| DB035621 | |

Abbreviations:

| D = Laboratory duplicate | |
|--------------------------|--|
| N = Native TCDD spike | |
| MB = Method blank | |

PE = Performance Evaluation Sample

RR = Rerun

NA = Non applicable

C = Critical error
M = Major error
m = Minor error
G = General error

* = See Interpretation Notes on p.8

Contractual Comments:

MINOR ERRORS:

- ml The batch number was incorrect on all forms. The correct batch numbers should be DB0355 and DB0356.

The formula used by the laboratory had "N" in the denominator and not "N-1".

General Comments:

- G1 Sample DB035524 is a Region 2 PE sample. The spike concentration and acceptance windows are not available to the data evaluator.
- G2 Sample results for DB035523 were not submitted because the internal standard recovery was zero. This sample is being reextracted and analyzed in another QC batch as stated in the case narrative. See Enclosure 1.
- G3 The analysis date for the initial calibration was 2/6/87. The samples were analyzed 10 months later on 12/4 and 12/5/87.
- G4 The laboratory did not follow the EPA rounding rules to report the mean RRF values on Form B-2.
- G5 The laboratory was inconsistent in choosing surrogate peak areas between the initial calibration data and sample data. See Enclosures 6 and 7.
- G6 The results for the percent recovery of the internal standard could not be duplicated. The internal standard recovery for samples MB, DB035515, DB035619 and DB035621 are below the advisory limit.
- G7 The percent recovery of the spiked field blank was miscalculated. The reported value, 98%, and the correct value, 99%, are within criteria.
 - G8 The RPD was not calculated because 2,3,7,8-TCDD was not detected in the sample, DB035614, and the duplicate sample, DB035614D.

ENCLOSURES:

- 1. Case Narrative
- 2. Form I
- 3. Form II
- 4. Form III
- 5. Form IV
- 6. Surrogate chromatogram for CCl analyzed on 2/6/87 at 01:10.
- 7. Surrogate chromatogram for DB035514

I. <u>DATA SUMMARY FORM</u> (Form B-1) (Exhibit B, Sec. A, p. B-11)

- *A. Form submitted for each matrix and all samples included on form
- B. Header information (Lab, Case/Batch No., instrument ID, report date, column)
- C. EPA sample number with proper suffixes
- D. Extraction date and GC/MS Analysis Date and Time
- E. Weight (to nearest 10th of a gram)/ volume (to nearest 10th of a ml)
- F. Calculated concentration of 2,3,7,8-TCDD (in correct units) if detected. Use 3 significant figures if >10 µg/kg or 100 ng/L and 2 if less than these quantities
- G. If TCDD not detected (ND), report a MPC (Ex D, 12.2, p.D-28)
- H. Signal to Noise (S/N) of Surrogate
- I. Recovery of the Internal Standard
- J. Raw peak areas for m/z 259, 320, 322, 328, 332IS, 334IS, 332RS and 334RS
- K. Relative Response Ratios for 320/322, 332/334IS and 332/334RS
- L. No calculation or typographical errors on the Data Report Form

II. <u>INITIAL CALIBRATION SUMMARY</u> (Form B-2) (Exhibit B, Sec. B, p B-15)

- *A. Form submitted for each instrument
- B. Header information (Lab, Case/Batch No., CC Solution Alternative, Instrument ID)
- C. GC/MS Analysis Date and Time
- D. Peak area for each ion: 259,320,322, 328, 332IS, 334IS, 332RS and 334RS
- E. Relative Response Ratios for 320/320, 332/334IS, and 332/334RS
- F. Relative Response Factors for the Native TCDD (RRFn) and the Internal Standard (RRFi)
- G. No calculation or typographical errors on Form B-3

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| III. | ROUT | INE | CALI | BRAT | CION | I SUMMARY | (Form | B-3) |
|------|------|-----|------|------|------|-----------|-------|------|
| | | _ | | | | B-18) | | |

- *A. Form submitted for each instrument and all PCS's and CC1's included on form
- B. Header Information (Lab, CC Solution Alternative, Case/Batch No, Instrument ID)
- C. GC/MS Analysis Date and Time
- D. Peak area for each ion: 259,320,322, 328, 332IS, 334IS, 332RS and 334RS
- E. Relative Response Ratios for 320/320, 332/334IS, and 332/334RS
- F. Relative Response Factors for the Native TCDD (RRFn) and the Internal Standard (RRFi)
- G. % Valley for PCS
- H. No calculation or typographical errors on Form B-3

IV. QUALITY CONTROL SUMMARY (Form B-4)

(Ex. B, Sec. D, p. B-20)

- *A. Form submitted for each batch
 - B. Header information (Lab, Case/Batch No., Instrument ID)
 - C. Sample numbers for fortified field blank and duplicate analysis
- D. Accuracy of fortified field blank spike
- E. Relative difference for the duplicate analysis
- F. No calculation or typographical errors on Form B-4

V. OTHER DELIVERABLES (Ex. B, Index, p B-6)

- A. Case Narrative provided
 - 1. Analytical problems addressed
 - 2. Documentation problems addressed
- B. All quantitation reports and SIM mass chromatograms for calibration solutions and performance check solutions (one m for each missing document
- C. All quantitation reports and SIM mass chromatograms for samples, including QA samples (one m for each missing document)
- D. Chain of Custody and in-house laboratory control documents

| Defect | | 7.7.1 | | Summany |
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| Type | NA_ | Yes | No | Summary Comment |
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VI. CALIBRATION QUALITY ASSURANCE CRITERIA

- A. Column Performance Check Solution (Ex. D, 9.2.6.1, p. D-17)
 - 1. Analyzed at proper frequency on all instruments
 - Valley ≤25% between 2,3,7,8-TCDD and all other TCDD isomers
 - Ratio of m/z 320 to m/z 322 between 0.67-0.9
 - Ratio of m/z 332IS to m/z 334IS between 0.67-0.9
- B. Initial Calibration (Ex. D, 9.2.6.2, p. D-17)
 - Standards at contract specified concentration ranges (Ex. D, 7.6, D-11-13)
 - 2. MS Sensitivity: S/N >2.5 for m/z 259,320,332, and 328 and S/N >10 for m/z 332 and 334
 - 3. Ratio of m/z 320 to m/z 322 between 0.67-0.9
 - 4. Ratio of m/z 332IS to m/z 334IS between 0.67-0.9
 - 5. Variation of the RRF for native 2,3,7,8-TCDD at each concentration not >10% RSD
 - *6. RSD <10% for the 4 mean RRF's for 13C12-2,3,7,8-TCDD
 - *7. RSD <10% for the 4 mean RRF's for native 2,3,7,8-TCDD
 - 8. Calculations performed correctly
- C. Routine Calibration (Ex. D, 9.3.3,
 - p. D-20)
 - 1. MS Sensitivity: S/N >2.5 for
 m/z 259,320,322 and 328 and S/N
 >10 for m/z 332 and 334
 - 2. Ratio of m/z 320 to m/z 322 between 0.67-0.9
 - 3. Ratio of m/z 332IS to m/z 334IS between 0.67-0.9
 - *4. Relative Response Factor for native 2,3,7,8-TCDD must be within 10% of mean value established by the initial calibration analysis.
 - 5. Calculations performed properly

| Defect | | - | | Summary |
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| VII. | IDENTIFICATION CRITERIA | (Ex. | D, | 11.6, |
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| | - D 061 | | | |

- p. D-26)
- A. Retention time of sample component within 3 seconds of the retention time of 13C12-2,3,7,8-TCDD (IS)
- B. Integrated ion currents for m/z 259, 320 and 322 maximize simultaneously
- C. MS Sensitivity: S/N >2.5 for m/z 259,320,322 and 328 and S/N >10 for m/z 332 and 334.
- D. Ratio of m/z 320 to m/z 322 between 0.67-0.9
- E. Ratio of m/z 332IS to m/z 334IS between 0.67-0.9
- F. Recovery of the internal standard within the advisory window of 40-120%
- G. Failure to report the concentration of any sample that meets all the criteria for positive identification
- H. If a positive sample is above the calibration range, 1 g reextracted
- If TCDD not detected, MPC calculated properly.
- J. If MPC > 1 µg/kl (soil) or 10 ug/L (water), sample reextracted and reanalyzed

VIII. NATIVE TCDD SPIKE RESULTS (Ex. E, 4.2.2.3 p. E-4)

- A. One sample spiked at 1.0ppb per batch of 24 or fewer sample
- B. Recovery of native TCDD within 60-140%. If not, rerun sample (Ex. C, 2.2.5 p. C-3)
- C. Recovery properly calculated
- D. Retention time of native TCDD within 3 seconds of the retention time of the internal standard

IX. <u>LABORATORY DUPLICATE ANALYSIS</u> (Ex. E, 5.11, p. E-4)

- A. One duplicate analysis per batch of 24 or fewer
- B. RPD for the analysis <50%

X. BLANK ANALYSIS RESULTS (Ex. E, 4.1,

- p. E-2)
- A. At least one method blank analyzed per batch and per matrix of 24 or fewer samples

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URIGINAL (Red)

- B. No contamination (i.e., no signal at m/z 259,320 or 322 >2% of m/z 332 within ±5 scans of the m/z 332 peak maximum)
- C. If contamination, associated positive samples reextracted and reanalyzed

| Defect Type | NA | Yes_ | No | Commen |
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XI. PE SAMPLE RESULTS

- A. Concentration within acceptance windows
- B. No false positives reported

Interpretation Notes:

- IA. M if Form B-1 is not provided (all other catagories in I are NA). m for each sample not included on Form B-1, but raw data included.
- IIA. If Form B-2 not provided, all other catagories in II are NA.
- IIIA. If Form B-3 not provided, all other catagories in III are NA.
- IVA. If Form B-4 not provided, all other catagories in IV are NA.
- VI B6. If % RSD \geq 20, error is C; if 10 to 20, error is M.
- VI B7. If % RSD \geq 20, error is C; if 10 to 20, error is M.
- VI C4. If % RSD \geq 20, error is C; if 10 to 20, error is M.

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State of Rem Jersen DEPARTMENT OF ENVIRONMENTAL PROTECTION DIVISION OF ENVIRONMENTAL QUALITY CN 027, TRENTON, N.J. 08625

JORGE H. BERKOWITZ, Ph.D. DIRECTOR

(609) 292-5383

MEMORANDUM

TO:

Andrew Fishman

Contract Administrator

Office of Quality Assurance

FROM:

Michael W. Miller, Ph.D.

Office of Quality Assurance

Floyd Genicola

Environmental Scientist I Office of Quality Assurance

SUBJECT: Audit Report, E.T.C. Corp. for Contract X-195

DATE:

February 23, 1988

On February 10, 1988 the above personnel conducted an on-site audit of E.T.C. Corp., Edison, New Jersey. We met with John E. Farre__, Technical Manager, Karen Kotz, QA Director, June Baker, QA Coordinator and laboratory personnel to discuss analytical methods, quality assurance and instrumentation. also reviewed a typical data package.

We recommend that E.T.C. Corp. performance for contract X-195 be rated conditionally accepted pending correction of data reporting deficiencies.

DEFICIENCIES

The Volatile Organic Analysis Blank reported with a sample (N.J.DEP, BC8488) is not the actual method blank analyzed previous to the sample set and within 12 hours. The blank reported in the data summary as the "method blank" is a screening blank analyzed a day earlier. The contract requires that the blank reported as the method blank be the actual blank analyzed with the sample. E.T.C. must report the actual method blank for every sample data summary.

AR304086

E.T.C. must identify all N.J.DEP VOA samples analyzed in the E.T.C. screening program. New data summary sheets must be issued to the N.J.DEP project manager with the correctly associated method blank for each sample.

- 2. Current procedures for mass spectral interpretation for Nontarget or Tentatively Identified Compounds are deficient. Data reported for sample BC8488 contained incorrect compound identifications. Contract guidelines for interpretation of mass spectra must be followed. E.T.C must improve the mass spectral identifications made by interpretation specialists.
- 3. Analysts in the Inorganics Section are not initialing data sheets. Data tabulation sheets for all inorganic methods are filled in by hand. The analyst must initial and date each sheet.
- 4. The organics preparation supervisor stated that Task IV is being cleaned by GPC. Gel Permeation Chromatography cannot be used for N.J.DEP Task IV acid/base-neutral extractables. Gel permeation is only acceptable for pesticides/PCBs.

RECOMMENDATIONS

- A. Changes in SOPs
 - 1. S-P-0-058: Disposal of Unused Samples

Samples are retained according to the customer's need or contract. Sample disposal shall be with written permission of the customers project officer.

- 2. SM-0-100: Log in Procedure
- a. N.J.DEP Chain of Custody Forms must be packed in every shuttle
- b. Shuttles must be shipped to the site unless arrangements are made for pick-up by the N.J.DEP project manager.
 - 3. Instrument and Method Detection Limits

A SOP for the determination of IDL's and MDL's is needed.

- 4. SOP's needed for the analysis of Petroleum Hydrocarbons in Solids and the analysis of TOX in Soils.
- 5. A complete set of corrected current SOP's should be sent to N.J.DEP-OQA by March 21.

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B. Laboratory Evaluation



- 1. Trip Blanks
- a. Trip blanks for organic analyses should be drawn from the same source as the Instrument Method Blanks.
- b. The trip blanks should be dated so that it is traceable to instrument blanks.
- 2. CLP reporting forms for VOA surrogates and aqueous matrix spike data are dated Rev 7/85 whereas BNs are dated Rev. 1/87. E.T.C. should correct <u>all</u> forms to the current CLP IFB.
- 3. E.T.C. must control the VOA Method Blank contamination to less than the CRDI. Methylene chloride must be controlled to less than 5 ppb. This will be enforced unless the contract is amended.
- 4. Currently, the Total Ion Chromatograms for extractables are presented as a 4" \times 4" chart. The complex TIC is very difficult to read. A two page presentation of the TIC is requested.

W JERSEY DEPARTMENT OF ENVIRONMENTAL PROTECTION OFFICE OF QUALITY ASSURANCE

PROFESSIONAL ANALYTICAL SERVICES CONTRACTS COMPLIANCE AUDIT (ON-SITE ADMINISTRATIVE AUDIT)

| Laboratory Name | Environmental Testing of 2017 the dien one |
|--|---|
| Address | Lingitar Center Parkery |
| <u></u> | 1 115 0 32/10-7 923 |
| Telephone No. | 1-22-5-6772 Date of Audit 1/1/19 |
| Document No. | SS - 025-2/88 Contract: X-195 XRI/FS Other: X |
| evaluated a N/A will authorized if required | A check mark will be made next to the item number to indicate a deficiency. If an item is not be placed in the remarks/comment area of the section not evaluated. Continuation sheets are d. All violations of N.J.A.C. 7:18 by a certified laboratory shall be repsorted to the Laboratory EP Office of Quality Assurance. |
| 7.8.1/B () Lab | oratory Operations were deficient in that: |
| types of analyses requ | aboratory did not have sufficient properly qualified personnel commensurate with the workload and tired to be performed pursuant to the Regulations Governing Laboratory Certification and Standards A.C. 7:18-2.7) or the most recent USEPA CLP IFB. |
| experience in laborate | e laboratory did not have a <u>Quality Assurance Officer (QAO)</u> with at least one (I) year ory quality assurance or quality control procedures and report directly to the laboratory manager wel of management in the same chain of command. |
| 7.8.2/B.3-10 () | Laboratory Personnel Requirements |
| 7.8.2.1/B.3 () | The GC Operator did not have at least nine (9) months experience in the operation of a GC on environment samples. N.J.A.C. 7:18-2.7(b)1 |
| REMARKS: | |
| | |
| 7.8.2.2/B.4 () | The GC/MS Operator did not complete a formal training course in GC/MS and have at least nine (9) months experience in the operation of the GC/MS data system on environmental samples. N.J.A.C. 7:18-2.7(b)2 |
| REMARKS: | |
| | |
| 7.8.2.3/B.5 () | The Extraction/Concentration Specialist did not have at least one (1) year experience in the preparation of extracts form environmental samples. N.J.A.C. 7:18-2.79b)3 |
| REMARKS: | |
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Form DEP-066 (11/87)

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| /.8.2.4/B.6 | (| , | purge and trap technique for volatile organic analysis. N.J.A.C. 7:18-2.7(b)4 |
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| REMARKS: | | | |
| 7.8.2.5/B.7 | (|) | The <u>Pesticide and Herbicide Residue Specialist</u> did not have at least two (2) years experience in Organochlorine/Organophosphorous pesticide, herbicide and PCB analysis including method specified cleanup procedures (i.e. column chromotography) on evironmental samples. N.J.A.C. 7:18-2.7(b)5 |
| REMARKS: | | | |
| 7.8.2.6/B.8 | (|) ~, | The Mass Spectral Interpretation Specialist did not have at least two (2) years experience in the interpretation of mass spectra generated from GC/MS analysis of environmental samples. N.J.A.C. 7:18-2.7(b)6 |
| REMARKS: | | | |
| 7.8.2.7/B9 | (|). | The Atomic Absorption Spectrometer Operator did not have at least six (6) months experience in the operation of atomic absorption equipment. N.J.A.C. 7:18-2.7(b)7 |
| REMARKS: | | | |
| 7.8.2.8/B10 | (|) | The <u>Inductively Coupled Plasma Operator</u> did not have at least nine (9) months experience in the operation of ICP equipment. N.J.A.C. 7:18-2.7(b)8 |
| REMARKS: | | | |
| 7.8.2.9 | (|) | The <u>Phase Contrast Microscopist</u> did not have at least one (1) year experience in the operation of a phase contrast microscope (PCM) or has not completed a formal training course in the operation of the PCM and associated equipment. |
| REMARKS: | | | |
| 7.9/C | (|) | Equipment Requirements were deficient in that: |
| 7.9.1/C.2 | |)a | The laboratory participating in Tasks I, II, III, V, VI, and VII or non USEPA CLP laboratory participating in RI/FS projects did not meet and maintain the minimum standards for laboratory nstrumentation set forth in the Regulation Governing Laboratory Certification and Standards of Performance, N.J.A.C. 7:18-1.1 et seq. |
| | (|)b | The laboratory did not meet and maintain the equipment requirements set forth in the analytical method bid. |

(OOA-LSS-008-11/87) ()c The laboratory did not maintain for archival storage of all data, except for GC/MS a bound, paginated and signature certified notebook. REMARKS: ()a The laboratory participating in Task IV or a CLP laboratory performing in RI/FS projects did not 7.9.2/C.1 meet and maintain the minimum standards for laboratory instrumentation set forth in the most recent USEPA CLP IFB document. ()b The laboratory did not maintain for archival storage of all data, except GC/MS, a bound paginated and signature certified notebook. REMARKS: () Additional Requirement for GC/MS Analyses-All Tasks, were deficient in 7.9.3/CLP that: () The mass spectrometer was not equipped with a computerized MS library search system capable of 7.9.3.1 providing reverse searching for targeted analytes and forward searching for non-targeted analytes. Software______ No. of Library Entries_____ () For archival storage of all GC/MS data the laboratory did not maintain a nine (9) track magnetic 7.9.3.2 tape system capable of archival storage of all data obtained in a form that can be retrieved on line to the data system. () The storage medium was not maintained under secure and appropriate conditions to preclude to 7.9.3.3 prevent loss of data. REMARKS: 7.9.3.4 () A permanent service record was not maintained in a logbook for each analytical instrument and ancillary equipment. REMARKS: 7.9.3.5 () An analytical instrument has been modified in an unacceptable manner. REMARKS: 7.9.3.6 () An analytical instrument was not adequately or properly vented.

REMARKS:

| 7.9.3.10 | (|) | Calibration intensity and gains were not kept in a permanent logbook for all calibrated instruments. |
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| REMARKS: | | | |
| 7.9.3.11 | (|) | Analytical balances were not calibrated within one (I) year by a certified technician. N.J.A.C. 7:18-4.7(e)I |
| REMARKS: | | | |
| 7.9.3.12 | (|) | Hood(s) were not in functional condition, flow rate monitored, and recorded in a logbook as required by N.J.A.C. 7:18-4.2(d). |
| REMARKS: | • | • | |
| 7.9.3.13 | (|) | The conductivity/res. tivity of distilled or demineralized laboratory water was not routinely checked and recorded in a permanent logbook. NJ.A.C. 7:18-4.6(b) |
| REMARKS: | | | |
| 7.9.3.14 | (|) | Analytical balances were not checked daily with the appropriate range of class S weights and the results recorded in a permanent logbook. N.J.A.C. 7:18-4.6(k) |
| REMARKS: | | | |
| 7.9.3.15 REMARKS: | (|) | The instrument manufacturer's operating manual was not readily available to the operator. |
| 7.9.3.16 | (|) | The laboratory cannot document any preventative maintenance program (internal or contracted) for analytical instruments and allied equipment. |
| REMARKS: | | | |
| 7.10/CLP | (| • | Sample Handling was deficient in that: |
| 7.10.1 | (|) | The appropriate portion of the laboratory SOP was unavailable to the sample custodian in the sample receipt area. $AR304092$ |
| 7.10.2 | (|) | The appropriate portion of the laboratory SOP was unavailable to the analyst. |

| (OQA-LSS-008 | -11/ | /87) | |
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| 7.10.3 | (|) | The employees of the laboratory were not following the laboratory SOP as written. |
| REMARKS: | | | $\frac{Q_{r_{i}}}{r_{i}}t_{r_{i}}$ |
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| 7.10.4 | (|) | Sample shipping containers were opened in a manner which did not prevent possible contamination of the laboratory or other samples. |
| REMARKS: | | | |
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| 7.10.5 | (|) | Aqueous samples (Tasks II and VI) were not preserved in accordance with the most recent 40 CFR 136 (water/wastewater) or 40 CFR 141 (drinking water). |
| REMARKS: | ; | | |
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| 7.10.6 | (|) | Samples collected and submitted under Task IV or submitted as part of a RI/FS project did no comply with the sample holding and preservation requirements of the most recent USEPA CLI IFB document. |
| REMARKS: | | | , |
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| 7.10.7 | (|) | Non-aqueous soil, sediment, and sludge samples (Non-CERCLA, Tasks III and IV) were not stored at 4.0 degrees C. |
| REMARKS: | | | |
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| 7.10.8 | (|) | Adequate facilities were not provided for the storage of samples. |
| REMARKS: | | | - 0k |
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| 7.10.9 | (|) | The temperature of the cold storage areas was not monitored daily and recorded in a permaner logbook. N.J.A.C. 7:18-4.7(e)6 |
| 7.10.11 | (|) | Temperature excursions (+/-4.0 deg. C) were noted. No corrective action was indicated. |
| 7.10.12 | (|) | The sample receipt/temperature records were not maintained in a appropriate manner. |
| REMARKS: | | | |
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| 7.10.13 | (|) | The laboratory was not maintained in a clean and organized manner. |
| 7.10.14 | (| ١ | Contamination free areas were not provided for trace level analytical work. |

() Contamination free areas were not provided for trace level analytical work.

7.10.14

| (OQA-LSS-008 | 8-11/87) | | 20. |
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| 7.10.15 | () | Reference materials were not labeled with concentrations, date of prepara the individual who prepared references or were not traceable in a perman standards were not stored separately from samples. | tion, and the identity of ent logbook. Reference |
| 7.10.16 | () | The laboratory did not possess a limited access, chemically isolated area for as dioxin or mixed waste. | r high hazard work such |
| 7.10.17 | () | The chemical waste disposal policies/procedures are not being followed or a | re inadequate. |
| REMARKS: | | <u>CK</u> | |
| 7.12.2 | () | Requirements for Aqueous Sample Analysis | |
| 7.12.2.1 | () | Purgeable Organics by GC (EPA 601, 602, and 603) is deficient in that: | |
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| 7.12.2.2 | X | Purgeable Organics by GC/MS (EPA 624) is deficient in that: | 15 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 |
| 7.12.2.3 | () | Extractable Organics (except pesticides and PCBs) by GC (EPA 604, 6 612) was deficient in that: | 607, 609, 610, 611, and |
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| 7.12.2.4 | () | Extractable Organics by GC/MS (EPA 624) was deficient in that: | |
| | | | |
| 7.12.2.5 | () | Pesticide and PCB Analysis (EPA 608) was deficient in that: | : |
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| 7.12.2.6 | () | 2,3,7,8-Tetrachlorodibenzo-p-dioxin (EPA 613 Analysis and/or 625 Screen |) was deficient in that: |
| | | | 4R304094 |

| 7.12.2.7 | \bowtie | Metal Analysis by Flame AA and/or ICP was deficient in that: |
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| | | initials of for a filling out tal sace trup little will by on late sheets. |
| 7.12.2.8 | (×) | Metal Analysis by Furnace AA was deficient in that: |
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| 7.12.3 | () | Requirements for Non-Aqueous Samples |
| 7.12.3.1 | (,) | Purgeable Organics by GC (SW-846 Methods 8010, 8020, and 8030) were deficient in that: |
| | | |
| 7.12.3.3 | () | Extractable Organics by GC except Pesticides and PCBs (SW-846, 8040, 8060, 8090, 8100, and 8120) were deficient in that: |
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| 7.12.3.4 | () | Extractable Organics by GC/MS except pesticides and PCBs (SW-846 8250 and 8270) were deficient in that: |
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| 7.12.3.5 | () | Pesticide/PCB Analysis by GC (SW-846 8080) was deficient in that: |
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| 7.12.3.6 | () | Polychlorinated Dibenzodioxins and Polychlorinated Dibenzofurans (SW-846 8280) was deficient in that: |
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| 7.12.6.2 | (|) | Purgeable Organics by GC/MS (EPA 624 Modified) was deficient in that: |
|----------|---|---|--|
| 7.12.6.3 | (|) | Extractable Organics (except pesticides and PCBs) by GC (EPA 604, 606, 607, 609, 610, 611, and 612) were deficient in that: |
| 7.12.6.4 | (|) | Extractable Organics by GC/MS (EPA 624 Modified) was deficient in that: |
| 7.12.6.5 | (|) | Pesticide and PCB Analysis (EPA 608 Modified) was deficient in that: |
| 7.12.6.6 | (|) | 2,3,7,8-Tetrachlorodibenzo-p-dioxin (EPA 613 and 625) were deficient in that the requirements set forth in Section 7.12.6.6 of the RFP were not met. |
| 7.12.6.7 | (|) | Polychlorinated dibenzo-p-dioxins and polychlorinated dibenzofurans (C/4 through C/8 congeners, SW-846 Method 8280, 40 CFR 261, Appendix X, 6 October 86) was deficient in that: |
| 7.12.6.8 | (|) | Metal analysis by Flame AA and ICP was deficient in that: |
| 7.12.6.9 | (|) | Metal analysis by Furnace AA was deficient in that: |
| | | | AR304097 |

| 7.12.7 | () | Requirements for Non-aqueous Samples (Task VII) |
|----------|-----|--|
| 7.12.7.1 | () | Purgeable Organics by GC (SW-846 8010, 8020, and 8030) were deficient in that: |
| 7.12.7.2 | () | Purgeable Organics by GC/MS (SW-846 8240 Modified) were deficient in that: |
| 7.12.7.3 | ; | Extractable Organics by GC (except Pesticides and PCBs), SW-846 8040, 8060, 8090, 8100, and 8120) were deficient in that: |
| 7.12.7.4 | () | Extractable Organics by GC/MS (except Pesticides and PCBs), SW-846 8250 and 8270 Modified) were deficient in that: |
| 7.12.7.5 | () | Pesticide/PCB Analysis (SW-846 8080 Modified) was deficient in that: |
| 7.12.7.6 | () | 2,3,7,8-Tetrachlorodibenzo-p-dioxin Analysis USEPA CLP IFB WA84-A002, 12/30/83 or the latest revision was deficient in that: |
| 7.12.7.7 | () | Polychlorinated Dibenzo-p-dioxins ans polychlorinated Dibenzofurans (C1/4 through C1/8 congeners), SW-846 Method 8280, 40 CFR 261, Appendix X, 6 October 86 was deficient in that: |

Name (Print):

Signature:

Title:

(OQA-LSS-008-11/87)

ON-SITE LABORATORY EVALUATION

LABORATORY PERSONNEL

| 20/2 | |
|--------|--|
| 3/1/4 | |
| - 52 \ | |

| DPESS | | ر کسی مسالات ا | PHONI | DN |
|----------------|---|-------------------|--|------------------------|
| 7,72,17,00 | | | | |
| NAME AND TITLE | EDUCA DEGREE PhD, MS, BS, BA,Assoc.,HS | MAJOR | NO. OF YEARS EXPERIENCE IN ENVIRONMENTAL TESTING | PRIMARY RESPONSIBILITY |
| Ku-1 -====== | | <u> </u> | | 141731 |
| | | | | Substitute of the |
| | | | | 1 11.52 |
| · 218 1 21/1 | | | | " E.L. / . = " |
| [) 宣言。三人 | | | | St. Contritt |
| | | ,· · . | | |
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| | | | | |
| | | | | |
| | | | | AR3041 |

Manager's Signature ____

NOITALLAND YNOTANCEAL STIENO

LABORATORY EQUIPMENT

| TYPE OF EQUIPMENT | YES | ABLE NO | MANUFACTURER | MODEL | SERIAL # | COMMENTS |
|------------------------------------|--|-------------|-----------------|-------------|--|-------------|
| | | . 1 | | | | |
| SERVICES | | | | | | CSEP |
| Light | | 1 | | | | |
| Electrica' | · _ | i | | | | |
| Gas | · | | | | | |
| Centra' Vacuum | | | | | 1 | |
| Secured Space | ئـــــــــــــــــــــــــــــــــــــ | | · | | | |
| Air Conditioning | | | | | | |
| LABORATORY WATER SUPPLY: | | } | | | | CLV+T |
| Distilled | , | | | | | |
| Double Distilled | | | | | | |
| Deionized | f | | | | | |
| | | | | | | |
| CHEMICAL STORAGE: | | | | | | 0070 |
| Volatile, Carcinogenic & Fiammable | | | | | | <u> </u> |
| Acids | | | | | | |
| | - | | | | - | OLION A. |
| Housekeeping | 1 | | | | | CHOK 0. |
| | | | | | | |
| EQUIPMENT: | | | | | | CVGL |
| Glassware (Class A volumetric) | - | | | | | |
| Pipets | - | | | | | |
| Burets | | | | | | |
| Flasks | | | | | | |
| Analytica! Balance | V | | LUERICUI SOFT | 155180 | 20,074 | |
| Pan Balance | + | | | | | |
| Top Loading Balance | | | , | | | |
| D.O. Meter | | | | | | |
| pH Meter | | | | | | |
| Buffer pH4 pH7 pH10 | | | | | | |
| Specific Ion Meter | | | | | | |
| Conductivity Meter | | | | | - | |
| Amperometric Unit | + | | 110011 1112 | V 0 | 00000 | |
| Turbidimeter | 1 | | HACH RATIC | XR | 870100232 | |
| Spectrophotometer (U.VVIS.) | | | | 1210 | 861101033 | O: D-OK-1 |
| Spectrophotometer (I.R.) | V | | YE | 10.0 | 707494 | |
| Filter Photometer | + | | | | | |
| Flame Photometer | +/ | | | | | |
| Mercury Analyzer | +-/- | | 16 C. 1811 . A. | | | 11-4 : |
| Auto Analyzer | + | | TECHNICY T | | | allate, de |
| Class S Weights | + | | | <u> </u> | | |
| NBS Thermometer | | | | <u> </u> | | ALLOY. |
| Total Organic Carbon Analyzer | + | | | | | CWGT |
| IR Detector | + | | | | | CTHM |
| FID Detector | - | | | | | |
| TOX Analyzer | | | | | | AR304101 |
| BOD Incubator | + | | | | | 1011444114 |
| Ion Chromatograph | | | | <u> </u> | | |
| Microbio Incubator | | | | | | |
| 44.5°C. Waterbath | | | | | | |
| Autoclave | | | | | | |

ON-SITE LABORATORY EVALUATION



LABORATORY EQUIPMENT (continued)

| TYPE OF EQUIPMENT | AVAILABLE YES NO | MANUFACTURER | MODEL | SERIAL # | COMMENTS |
|--|--|--------------|--|--|-------------|
| 170°C Quen | | - | | _ | |
| Aram : Absorption | | | | | |
| Piasma Spectrometer | , | | | | 2.5 |
| DCP | | | | | |
| ICAP | | | | | |
| Gas Chromatograph | | | | | |
| Refrigerator | | | | | |
| Freezer | | , | | | |
| Drying Oven | | L 12 + 4 5: | サントスト | 17:17 (3) | |
| Muffle Furnace | | | - <u>!</u> - | | |
| Hot Plates | | | | | |
| Magnetic Stirrer | | | | | |
| Desiccators | , | | | | |
| Steam Bath | | | | | |
| Stirred Boiling Water Bath with | | | ı | , | |
| Gabled Lid for Nitrate by | | | i | | |
| Brucine Method | · ; | | | ļ. | |
| Centrifuge | | | _ | 1 | |
| LABORATORY APPARATUS: | | | | | - |
| Fluoride Distillation | | , | : | : | |
| COD Reflux | , | - 1 4 4 1 | v. 3.4 | 100 | |
| Kjeldahl | i | | | | |
| ieldahl Digester | | | | 1 , | |
| anide Distillation | | | | 1 | |
| Soxhlet Extraction | | , | | | |
| LABORATORY SAFETY: | | | | | |
| Emergency Exits | | | | i | |
| Fire Alarm | , | | Ī | | |
| Smoke Detector | 1 | | 1 1 | | - VIV on |
| Sprinkler System | | <u> </u> | | | |
| Fire Extinguishers | <u> </u> | | i | | |
| Fire Blanket | | | | 1 | |
| Emergency Lights | - | | | | |
| First Aid Station | | | <u> </u> | | |
| Emergency Phone Numbers | | | | | |
| Hazardous Materials Chart | | | | <u> </u> | |
| Eye Wash Stations | | | | | |
| Chemical Burn Stations | | | | | |
| Safety Shower | | | | | |
| Lab Coats | | | | | |
| Safety Glasses | | | 1 | | |
| Face Shield | | | + | | |
| | - | | | 1 | |
| Respirator with Compressed Air Supply | | | | | |
| Fume Hoods | i i | | | | |
| Perchloric Acid Hood | | | | <u> </u> | |
| Compressed Gas Tanks Secured | | | <u> </u> | | |
| lectrical Cables Secured | | ; | | 1 , | AR304102 |
| s there an antidote for HF burns? e.g. A paste of MgOH and Glycero and a saturated solution of MgSO4 | 4 | | | | |

ON SITE LABORATORY EVALUATION

LIMITED CHEMISTRY GENERAL PROCEDURES

Ohi.

| | <u>V.4</u> |) ES | <u>NO</u> | <u>COMMENTS</u> |
|---|------------|-------|-------------------------------------|-----------------|
| ACIDITY | _ | | | |
| Are sample containers field completely? Are samples analyzed within 14 days of collection? Is the NaOH titrant standard zed against potassium. | | | CACD 01 | A |
| biphthalate and labeled properly? | | _ | CACD 03 | |
| 4 Are wastewater sample, titrated to pH 8.3 using an electrometric endpoint? | | _ | CACD 04 | |
| 5 If a phenophthale-in indicator is used, is free residual chlorine removed with thiosulfate? | | | CACD 05 | |
| ALKALINITY | | , | | |
| 1. Are sample containers filled completely? 2. Are samples analyzed within 14 days of collection? | | FIE | CALK 01 | |
| 3 Is the HaSO4 or HC1 standard to against Nac CO3 and labeled protective. | | | CALK 03 | |
| Are wastewater samples titrated to pH 4.5 using an electrometric endpoint? | | · | /□ CALK 04 | |
| 5. If methyl orange indicator is used, is free residual chlorine removed with thiosulfate? | | T | CALK 05 | |
| BIOCHEMICAL OXYGEN DEMAND | | | | |
| Are samples cooled to 4°C during transit and received in lab within 48 hrs, of collection? | | | CBOD 01 | |
| 2. Is the sodium thiosulfate standardized against potassium biniodate or potassium dichromate and labeled properly?3. Is a seed used on chlorinated or industrial effluents? | | | CBOD 02 | |
| 4. Is the depletion of unseeded dilution water blank less than 0.2 mg/1? | | | ☐ CBOD 04 | |
| 5. Do the sample dilutions used to compute the BOD have depletions of at least 2 mg/l and a residual DO of 1 mg/l? | | | ☐ C80D 05 | |
| 6. Is a glucose-glutamic acid standard included with approximately every 20 analyses?7. Is the BOD incubator thermometer graduated in intervals | | | CBOD 06 | |
| of 1 ^O C or smaller? | | | ☐ CBOD 07 | |
| 8. Is chlorine removed with sodium sulfite?9. How many dilutions are prepared to determine BOD? | ليا | | C80D 08 | 1 2 3 4 5 |
| CHEMICAL OXYGEN DEMAND (Holding 28 Days) | | | / | |
| Are samples preserved with H₂SO₄ to a pH to 2? Upon receipt in the laboratory, is the sample pH measured | | | CCOD 01 | |
| and recorded to verify that it is preserved? 3. Is the Dichromate reflux method used? | | Ø | CCOD 02 | |
| a. Is the ferrous ammonium sulfate titrant standardized daily against primary standard grade K₂Cr₂O₇? b. Is 0.025 N K₂Cr₂O₇ used for samples below 50 mg/l? c. Is a blank run with each set of samples? | | | ☐ CCOD 03 ☐ CCOD 04 ☐ CCOD 05 | |
| d. Is at least 0.5 ml of titrant used in the titratic of the excess dichromate for the majority of samples e. Is HgSO4 used to complex chloride? 4. Is the automated colorimetric method used? 5. Is the manual colorimetric method used? | | | CCOD 06 CCOD 07 CCOD 08 | AR304103 |
| a. Are digestion tubes heated in a block heater or oven at 150°C for 2 hrs.? | | D | | |

ON-SITE LABORATORY EVALUATION



| MARDNESS, TOTAL (Holding 6 mos.) | <u>NA</u> | YES | <u>NO</u> | | COMMENTS |
|---|-----------|-----|-----------|-------------------------|----------|
| 1. Are samples preserved with acid (HNO ₃ or H ₂ SO ₄) to pH<2? 2. Is the EDTA titrimetric method used? | | | | CHRD 01 | |
| a. Is the EDTA titrant standardized against CaCO3 and labeled properly? b. Is the EDTA titrant approximately 0.01M? 3. Is the automated colorimetric (calmagite) method used? | | | | CHRD 02 CHRD 03 | |
| 4. Is the hardness calculated from Ca+Mg values determined by atomic absorption? | | | | | |
| HYDROGEN ION (pH) (Analyze immediately) | | | | | |
| Is an electronic pH meter with temperature compensation used? Are electrodes stored according to the manufacturer's | | | | СрН 01 | |
| recommendations? | | | | CpH 02 | |
| 3. Are the electrodes filled with sufficient quantity of electrolyte? | | | | CpH 03 | |
| CONDUCTIVITY (Holding 28 Days) | | | | | |
| Are samples measured at 25°C or is a temperature correction made? Has the cell constant of the conductance cell been | | | | CCON 01 | |
| determined and permanently recorded? | | | | CCON 02 | |
| THYLENE BLUE ACTIVE SUBSTANCES solding 48 Hrs.) | | | | | |
| Is MBAS being determined by the methylene blue method? Is LAS reference material available and used in the | | | | CMBA 01 | |
| preparation of standards?3. Is the determination of absorbance done at 652 nm against a blank of chloroform? | | | | CMBA 02 CMBA 03 | |
| TURBIDITY (Holding 48 Hrs.) | | | | | |
| Is the nephelometric method used? Are samples with turbidity greater than 40 NTU diluted with turbidity-free water? | | ø, | | CTUR 01 | |
| b. Are sample tubes clear, colorless glass which are clean and have no scratches? | | 团 | | CTUR 02 | |
| COLOR (Holding 48 Hrs.) | | | | | |
| 1. Is the visual comparison method used? a. Is interference due to turbidity removed by filtration or centrifugation? b. Is the pH of the sample measured and reported with the result? c. Are platinum-cobalt standards used? | | | | CCOL 01 CCOL 02 CCOL 03 | |
| d. Are color disc stanceards described against platinum-cobalt standards every 6 months? Is the spectrophotometric method used? Is the ADMI method used? | | | | CCOL 04 | AR304104 |

ON-SITE LABORATORY EVALUATION

| RESIDUE, (T.D.S.), (TOTAL FILTERABLE RESIDUE) | <u>N 1</u> | <u>YES</u> | <u>NO</u> | COMMENTS |
|--|------------|------------------|--|----------|
| (Holding 48 Hrs.) Does the desiccator have suitable dessicant and indicator? Is an analytical balance capable of weighing to 0.1 mg available? Are glass fiber filter discs used? Are samples for total dissolved solids dried at 180°C? a. Does the dissolved residue, when weighed, yield <200 mg? b. If not, is smaller aliquot used? | | मुस्तिष्ठात् । १ | CTDS 01 CTDS 02 CTDS 03 CTDS 04 CTDS 05 CTDS 06 | |
| RESIDUE, (TSS), (TOTAL NONFILTERABLE RESIDUE) (Holding 7 Days) | | , | / | |
| 1. Is the residue dried at 103-105°C? | | | CTSS 01 | - |
| RESIDUE, TOTAL SOLIDS (Holding 7 Days) | | | , | |
| 1. Is sample dried at 103-105°C until weight is constant? | | Œ | | |
| CHLORIDE (No Pres., Holding 28 Days) | | | | |
| Is the argentometric (silver nitrate) method used? a. Is the AgNO3 titrant standardized against NaC1 dried at 140°C and labeled properly? b. Is interference due to sulfide, sulfite or thiosulfate | | | CCLD 01 | |
| removed with H2O2? | | | CCLD 02 | |
| 2. Is the mercuric nitrate method used? a. Is the pH adjusted to 2.5? b. Is a 1 or 5 ml microburet used for titration? c. Is the NaCL standard dried at 600°C for 1 hour? 3. Is the automated ferricyanide method used? 4. Is the ion chromatographic method used for drinking water? | | | CCLD 03 CCLD 04 CCLD 05 | |
| FLUORIDE (No Pres., Holding 28 Days) 1. Are water samples distilled? | | | ☐ CFLR 01 | |
| 2. Is the specific ion electrode method used? a. Are both samples and standards analyzed at room temperature? 3. Is the SPADNS method used? | | | ☐ CFLR 02 | |
| a. Is the SPADNS solution stored in an amber bottle and protected from direct sunlight? b. Is sodium arsenite used to remove residual chlorine? 4. Is the automated complexone method used? a. Is the working color reagent prepared fresh every | |] [] | CFLR 03 | |
| 3 or 4 days? | لا | لـا | CFLR 05 | |
| CHLORINE RESIDUAL (No Pres., No Holding) | | | | |
| Is chlorine residual determined by iodometric titration, DPD colorimetric or DPD titrimetric method: b. In the iodometric titration is the excess reducing agent | | | CCLR 01 | |
| back-titrated with iodine or iodate solutions | \Box | | CCLR 02 | 1000: |
| c. In the DPD colorimetric method are kits with polor | | \Box | | AR304105 |
| wheels, and reagent packets used? d. Is the chlorine residual determined by specific ion electrode? e. Is the starch end-point method used? |] [] | | | |

ON-SITE LABORATORY PROCEDURES



| | NA | YES | NO. | COMMENTS |
|---|-------------|--------------------------|-----------|--------------|
| SULFATE - Continued | | 123 | <u></u> | CO.M.HE. VIS |
| 2 b. Are both samples and standards read at 5 ± 0.5 | | · | | •• |
| minutes after stirring? | <u>`</u> _; | = | CSFA CSFA | |
| c. Are blanks used to correct for color or turbidity?3. Is the automated chloranilate method used? | | نے | L CSFA | 06 |
| a. Are interferences due to Ca, AI, and Fe removed by | | | | |
| an ion exchange column? | | 区/ | CSFA | 07 |
| 4. Is the ion chromatography method used for drinking water? | | $\overline{\mathcal{C}}$ | | |
| SULFIDE (Pres 4°C,Zn Acetate + NaOH to pH>9 - Holding 7 Days) | | | | |
| 1. Is the Methylene Blue method used? | | | | |
| a. Is the methylene blue solution standardized against a know | /n | | | |
| solution and adjusted so that 1 drop = 1.0 mg/l sultide? | | | ☐ CSFD | |
| b. Is the titrimetric (lodine) method used? | | | CSFD | 02 |
| SULFITE (No Preservation) | | | | |
| 1. Is the titrimetric iodine-iodate method used? | | | | |
| 2. Are samples analyzed on site? | | | ☐ CSFT | 01 |
| CYANIDE | | | | |
| Are samples analyzed within 14 days of collection? | | | ☐ CCYN | 01 |
| 2. Are samples preserved with NaOH to pH 12 + 0.6 G | _ | | | |
| ascorbic acid? | | | CCYN | 02 |
| Jpon receipt in the laboratory, is the pH measured and | | | | |
| recorded? | | | CCYN | 99 |
| 4. If chlorinated, do you remove sulfide as Cd sulfide? | 님 | 닏 | CCYN | 04 |
| 5. Is a manual distillation with MgCl2 done? 6. Is the titrimetric method used? | 님 | \exists | CCYN | 05 |
| a. Is the AgNO3 standardized against NaCl and labeled | | ليسيا | | |
| properly? | | | CCYN | 06 |
| b. Is a blank run with each set of samples? | | | CCYN | |
| 7. Is the colorimetric method used? | - | | | |
| a. Is Chloramine T prepared weekly and stored in | | | | |
| refrigerator? | | | CCYN | 08 |
| b. Is the stock cyanide solution standardized weekly | | $\overline{}$ | | 00 |
| against AgNO ₃ ? | لــا | نا | ☐ CCYN | <u> </u> |
| OIL AND GREASE (Holding 28 Days) | | | , | |
| 1. Are samples collected in glass containers? | | | CONG | |
| 2. Are samples preserved with H ₂ SO ₄ to pH<2? | 님 | | / ` | 02 |
| 3. Is a liquid-liquid extraction with freon used? | 뭄 | | ☐ CONG | |
| 4 Is the oil and grease content determined gravimetrically? | ب | لكنا | L CONG | 04 |
| PHENOLS (Holding 28 Days) | | | | |
| 1. Are samples collected in glass containers? | | | CPHN | |
| 2. Are samples preserved with H ₂ SO ₄ to pH<2? | | | ☐ CPHN | 02 |
| 3. Upon receipt in the laboratory, is the pH measured and | | | ☐ CPHN | 02 |
| recorded? | H | H | ☐ CPHN | |
| Are samples analyzed within 28 days of collection? Is the colorimetric 4AAP method with distillation used? | H | | ☐ CPHN | V+ |
| 6. Is the colorimetric 4AAP method for halogenated phenois | | | 5 | |
| used or | | | | |
| Is U.S.E.P.A. Method 604 used? | | | ☐ CPHN | 06 |
| | | | • | |

Form DWR-156 5 85

NEW JERSEY DEPARTMENT OF ENVIRONMENTAL PROTECTION OFFICE OF SCIENCE AND RESEARCH ON-SITE LABORATORY EVALUATION

RECORD-KEEPING AND CALIBRATION PRACTICES

| R | ECORD-KEEPING | NA. | YES | NO | | COMMENIS |
|-----------|---|-----|------------|------------|-------------------------|-----------|
| 1 | Is the temperature of all B.O.D. incubators recorded daily? | | | / □ | CREC 01 | |
| 2 | Is the temperature of all drying ovens recorded daily? | | | | CREC 02 | |
| 3. | Is the temperature of all refrigerators recorded daily? | | | | CREC 03 | |
| 4. | Are the laboratory thermometers calibrated against an NBS traceable thermometer and documented? | | V | | CREC 04 | |
| 5 | Is the analytical balance checked monthly with two class S weights, one in the mg range, and one in the gram range, and the data recorded? | | Z / | | CREC 05 | |
| 6. | Is a record available of yearly service on the analytical balance? | | Ø | | CREC 06 | |
| 7. | Is the pH meter checked daily, or before use, by setting the meter to pH7 then measuring and recording pH's approximately 4 and 10? | | | | CREC 07 | |
| 8. | Is the conductivity of the distilled water supply (satisfactory is conductivity of 2.0-0.5 umho/cm. at 25°C .) checked daily and the data recorded? | | | | CREC 08 | |
| 9. | Is the conductivity meter calibrated daily against a 0.001 M KCl solution and the data recorded? | | | | CREC 09 | |
| 10. | Is the turbidimeter calibrated daily, or before use, with a 40 NTU formazin standard and the data recorded? | | Y | | CREC 10 | |
| 11. | Is the DO meter calibrated weekly against the Winkler method and the data recorded? | | | | CREC 11 | |
| <u>C/</u> | ALIBRATION PRACTICES | | | | | _ |
| 1. | Regarding calibration curves, are the following practices in use? a. Graph is labeled with parameter, date of calibration and the axes are properly identified as to absorbance or percent transmission and concentration units. b. Computer read-out for regression analysis lists parameter, date of calibration, equation of curve and correlation co-efficient. c. Results reported are within the range of the highest and lowest standard. | | | | CCAL 01 CCAL 02 CCAL 03 | |
| 2. | Regarding manual spectrophotometric calibration curves, are the following practices in use? a. A minimum of 5 standards and a blank, with 3 measurements at each point are used to generate the curve. b. A new curve is generated every 6 months. c. The working curve is checked daily or with each run by alternating a low and a high standard and the data are recorded. | | 00 0 | | CCAL 04 CCAL 05 | |
| 3. | Regarding calibration curves for auto-analyzer analyses, are the following practices in use? a. The baseline is set using appropriate reagents and distilled water and is checked at the end of the run. b. A minimum of 5 standards are used to generate the curve. | | | | CCAL 08 | |
| | c. A new curve is generated for each run. d. A marking standard is included with every 20 samples. e. The calibration curve is checked at the end of each run with a low and a high standard and the data are recorded. | | | | CCAL 10 CCAL 11 | *P201.107 |
| 4. | Regarding atomic absorption calibration curves, are the following practices in use? | ت | ن_ | لب | JUAL II | ##384107 |
| | a. Working standards are prepared fresh with each run. | | | | CCAL 12 | |
| | b. A minimum of 4 standards and a blank are used to generate a curve.c. A new curve is generated for each run. | | | | CCAL 13 | |

ON-SITE LABORATORY EVALUATION

QUALITY CONTROL AND DATA HANDLING

| | <u>NA</u> | <u>YES</u> | <u>NO</u> | <u>COMMENTS</u> |
|--|-----------|------------|-------------------------|-----------------|
| QUALITY CONTROL | | | | |
| Regarding standard solutions, are the following practices in use A notebook record is available describing the preparation and standardization of stock standard solutions. Are purchased standards checked before use? | | | CQCS 01 | |
| c. Stock standard solutions and working standards are labeled with reagent, concentration, date prepared and initialed. d. ACS grade or analytical reagent grade chemicals dated when received, are used in the preparation of standard | | | □ cocs 03 | |
| solutions. | | | CQCS 04 | |
| Regarding the monitoring of precision, are the following practices in use? a. Approximately 1 synthetic known control sample is include with every 20 analyses, and the data presented on an X bar | đ | | | |
| control chart. b. Approximately 1 duplicate of a natural sample is included with every 20 analyses, and the data presented on an R | | | □ CQCP 01 | |
| bar range control chart. 3. Regarding the determination of chemical recovery, are the | لسا | | CQCP 02 | |
| following practices in use? a. A tabulation and control chart are available for recovery data obtained from spiked natural samples (1 for every | | | COCP 03 | |
| 20 analyses). 4. Is there an in-house quality control manual outlining Q.C. practices? | | | ☐ CQCP 04 | |
| DATA HANDLING | | | | |
| Regarding sampling procedures and data handling, data reporting and data retrieval procedures, are the following practices in use? | | | | |
| a. Are sample collectors supplied with properly labeled containers, preservatives and sampling instructions? (Get copy of instructions). b. Is there a lab daily work sheet listing sample number, | | | CDAT 01 | |
| date, time, location, preservation, analyses requested, field measurements by sampler, sampler's initials, date and hour received by lab, analysis, date and hour of analysis, analyst's initials? | | | CDAT 02 | |
| c. Is there a bound lab notebook for recording raw data, calculations, or other notes.d. Is raw data kept for 5 years?e. Is enforcement data kept for 5 years? | | सत्व | CDAT 04 CDAT 05 | use data skuzi. |
| f. Is there an in-house methods manual available to all analysts?g. Is there a record of chain of custody?h. Is there a chain of custody procedure? | | 000 | CDAT 06 CDAT 07 CDAT 08 | AR304108 |

ON-SITE LABORATORY EVALUATION

LABORATORY PERSONNEL

| LABORATORY ENVIRONMENTAL TESTINGS EXT | DATE OF | |
|---------------------------------------|----------------|--|
| ADDRESS 284 NACITAN GENTER | PHONE 525 674' | |
| 5-50N. N. T. 09837. | | |

| | EDUC | NO. OF YEARS | | | |
|------------------|--|----------------|-------------------------------------|----------------------------|--|
| NAME AND TITLE | DEGREE PhD, MS, BS, BA,Assoc.,HS | MAJOR | EXPERIENCE IN ENVIRONMENTAL TESTING | PRIMARY RESPONSIBILITY | |
| Tuie Edinfer | B 5. | pare. | 4 | SUPERUSIR GA. SUP. CER FLA | |
| 15. Klain | N. S. | 51. | / | GAT RESTAN | |
| Echn Fisz garaid | M.D. | Aun. Cheine | | NIPNAEEK. | |
| Kusen Kotz Espal | <u>,, / < .</u> | ۲., | | 5 12 1 5 A 4 A GC | |
| Tur defines | F. S. | | ٠. د | E reverse Behis | |
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| A | , | | | | |
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| 12/ - | | | | | |
| _33bec | | <u> </u> | | AR304109 | |

| Total ft ² of lab space 33600 | `. |
|--|-----------------------------------|
| Total linear feet of lab bench | Inspected by There was a superior |

ON-SITE LABORATORY EVALUATION





| TYPE OF EQUIPMENT | AVAII YES | ABLE NO | MANUFACTURER | MODEL | SERIAL# | COMMENTS |
|--------------------------------|--|--|--------------------|--|------------------|---|
| SERVICES: | | | | | | CSER - |
| Light | 1/ | | | | | 32: 2 |
| Electrical | 1 | /. | | 1 | 1 | |
| Gas | 1 | 1 | | | | |
| Vacuum | 11 | V | 1 | | 4 | |
| Secured Space | 11/ | | 2 / - 11/4 (1) 11 | 12/01/1 | 120814 | th |
| Air Conditioning | V_ | | | <i>II</i> | (9) | |
| LABORATORY WATER SUPPLY: | | | (| | | CLWT 🗆 |
| Distilled | | | | | | · |
| Double Distilled | | | | | / | |
| Deionized | | | guil y ter in | MED LINE | 11/1/11/11 | C I May |
| CHEMICAL STORAGE: | 1 | | | | | сѕто 🗆 |
| Volatile Reagents | 11/ | | | | | 33.0 |
| Acids | 10 | | | | | |
| Carcinogenic Reagents | 1/ | | | | <u> </u> | |
| Flammable Reagents | 1/ | | | | | |
| EQUIPMENT: | | | | | | CVGL 🗆 |
| Glassware (Class A volumetric) | | | | | <u> </u> | |
| Pipets | V | | | | 1 | |
| Burets | 10/ | | Print was lived | | | |
| Flasks | 1 | <u> </u> | | | | |
| Analytical Balance | V. | L)A | 15 TALES | L. 160 | 1. 3.19 | 124146 60 7 15/81. |
| Pan Balance | V , | 1 | 0.44.18 : | ļ | | |
| Top Loading Balance | | | ALMAN INTERNATION | d WHILE | | Vottles 7/25/56 |
| D.O. Meter | | V | | | | <u> </u> |
| ب H Meter | V | | ORILN | \$ | T | |
| Buffer | | | | <u> </u> | | |
| pH 4 | 17 | | | | | |
| pH 7 | V | //_ | | | | |
| pH 10 | 1/ | _/_ | reads (1.73 | <u> </u> | <u> </u> | |
| Specific Ion Meter | 1/ | V / | · | <u> </u> | ļ | |
| Conductivity Meter | \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ | 1// | V 43: | 31 | 1:17 | 17 |
| Amperometric Unit | | V | | | | |
| Turbidimeter | | 1 1/ | • | Ceria | | 1.2 |
| Spectrophotometer | ↓ ✓ | // | 2 | CAE. 4. | 1214A(0) | 4-1-3 |
| Filter Photometer | | <u>\</u> | | | | |
| Flame Photometer | /- | 10 | | - | | -, |
| Mercury Analyzer | + ./- | | And the same | 447 | 1 | a to the state of |
| Auto Analyzer | Y | } | the second | | CUAL : UM | CWGT L |
| Class S Weights | + 🏏 | ļ. —— | " with your | | NB 57/88 | |
| NBS Thermometer | + 1/ | | AIDS - ZOTE IN A | | W/25/188 | CITIM |
| Total Organic Carbon Analyzer | + - / | | L D I D I WENT WIN | T 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 | 7, +1/ | |
| →R Detector FID Detector | | | | | + | |
| | +- | | m x de l'est | \ | + | 1000:::0 |
| TOX Analyzer BOD Incubator | + - | - | my denisch | 1-1-6-1 | 1 | AR304110 |
| Microbio Incubator | + | | | + | | |
| 44.5°C. Waterbath | + | | | | - - | |
| Autoclave | | | | | + | |
| / ~u (UCIBTE | 1 | 1 | l . | 1 | ī | |

ON-SITE LABORATORY EVALUATION

LABORATORY EQUIPMENT (continued)

| | AVAIL | ABLE | | | | | |
|---|---|--|--|--|--|----------|--|
| TYPE OF EQUIPMENT | YES | NO | MANUFACTURER | MODEL | SERIAL # | COMMENTS | |
| 170°C. Oven | | | | | | | |
| Atomic Absorption | ¥ 1 | | 3125000 | | | | |
| Plasma Spectrometer | 1 | | 12- 2000 16500 | | | | |
| DCP | | | | | | | |
| ICAP | 7 | | /^ | | | | |
| Gas Chromatograph | 1 | | 12) Hil: | , , | | | |
| Refrigerator | V | | 125 /71: (10 | (b. 42) | | | |
| Freezer | | | - | , | | | |
| Drying Oven | 7/ | | The True | 1:46 | | | |
| Muffle Furnace | V | | The state of the s | | | | |
| Hot Plates | 7 | - | | | | | |
| Magnetic Stirrer | V | | | | | | |
| Desiccators | \$7.70 | 1 | in white | | | | |
| Steam Bath | V | | | | | | |
| Stirred Boiling Water Bath with | | <u> </u> | | | | | |
| Gabled Lid for Nitrate by | | [| | | | | |
| Brucine Method | 1 | | | | 1 | • | |
| Centrifuge | 1 | | 15. し | | | | |
| | | | | | | | |
| LABORATORY APPARATUS: | | | | | | | |
| Fluoride Distillation | <u> </u> | | | | | | |
| COD Reflux |), | α | | | | | |
| Kjeldahl | λ. | A | | | | | |
| Kjeldahl Digester | Α. | A | | | | | |
| Cyanide Distillation | | V | | | | | |
| Soxhlet Extraction | V | | ist; | | | | |
| LABORATORY SAFETY: | , | } | | | | | |
| Emergency Exits | / | | | | | | |
| Fire Alarm | 1 7 | 1 | | | | | |
| Smoke Detector | \ \tag{\tau} | | | | | | |
| Sprinkler System | | / | | | | | |
| Fire Extinguishers | V. | - | † | | | | |
| Fire Blanket | 3. | - | | <u> </u> | | | |
| Emergency Lights | | | | | | | |
| First Aid Station | 1/ | | | | | | |
| Emergency Phone Numbers | 107 | | | T | 1 | | |
| Hazardous Materials Chart | 1 | <u> </u> | | | 1 | | |
| Eye Wash Stations | 11/ | | | } | | | |
| Chemical Burn Stations | V. | 1 | | f | | | |
| Safety Shower | | | | | | | |
| Lab Coats | - | 1 | <u> </u> | 1 | † | | |
| Safety Glasses | 1/ | | | | | | |
| Face Shield | , , | / | | | † | | |
| Respirator with Compressed | | + / | | | | | |
| Air Supply | / | V , | | | 1 | | |
| Fume Hoods | V, | 1 | <u> </u> | - | | | |
| Perchloric Acid Hood | / | 1 1 | | | | AR304111 | |
| Compressed Gas Tanks Secured | \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ | 1 | | | | | |
| Electrical Cables Secured | 1 | | | | | | |
| Is there an antidote for HF burns? | | 1 ! | | | | <u> </u> | |
| e.g. A paste of MgOH and Glycerol and a saturated solution of MgSO4 | | ·j | | | | | |

ON-SITE LABORATORY EVALUATION

LIMITED CHEMISTRY GENERAL PROCEDURES

| | | NA | <u>YES</u> | <u>NO</u> | COMMENTS |
|--|-------|-------------|------------|------------------|-----------|
| ACIDITY | 4 = | | *1 | w.· | |
| Are sample containers filled completely? Are samples analyzed within 14 days of collection? | | | | CACD 01 CACD 02 | |
| 3. Is the NaOH titrant standardized against potassium biphthalate and labeled properly? | | | | CACD 03 | |
| 4. Are wastewater samples titrated to pH 8.3 using an electrometric endpoint? | | | | CACD 04 | |
| 5. If a phenophthalein indicator is used, is free residual chlorine removed with thiosulfate? | | | | CACD 05 | |
| ALKALINITY | | | | | |
| Are sample containers filled completely? | | | \Box | CALK 01 | · |
| 2. Are samples analyzed within 14 days of collection? | | | \equiv | CALK 02 | |
| 3. Is the H ₂ SO ₄ or HCl standardized against Na ₂ CO ₃ an labeled properly? | d | | | CALK 03 | |
| 4. Are wastewater samples titrated to pH 4.5 using an electrometric endpoint? | | | \Box | CALK 04 | |
| 5. If methyl orange indicator is used, is free residual chlo removed with thiosulfate? | rine | | | | |
| BIOCHEMICAL OXYGEN DEMAND | | - | | | |
| 1. Are samples cooled to 4°C during transit and received | in | | | | |
| lab within 48 hrs. of collection? s the sodium thiosulfate standardized against potassiu | m | | | CBOD 01 | |
| iniodate or potassium dichromate and labeled proper | | | | CBOD 02 | |
| 3. Is a seed used on chlorinated or industrial effluents? | | | | CBOD 03 | |
| 4. Is the depletion of unseeded dilution water blank less than 0.2 mg/l? | | \Box | \Box | CBOD 04 | |
| 5. Do the sample dilutions used to compute the BOD have | | _ | | | |
| depletions of at least 2 mg/l and a residual DO of 1 mg 6. Is a glucose-glutamic acid standard included with approximately acid. | | | | CBOD 05 | |
| mately every 20 analyses? | 0.21. | | | CBOD 06 | |
| 7. Is the BOD incubator thermometer graduated in interv | /als | | .— | ☐ Ć8OD 07 | |
| of 1 ^O C or smaller? 8. Is chlorine removed with sodium sulfite? | - | | | ☐ CBOD 08 | |
| 9. How many dilutions are prepared to determine BOD? | | - | | | 1 2 3 4 5 |
| CHEMICAL OXYGEN DEMAND (Holding 28 Days) | | | | | |
| 1. Are samples preserved with H ₂ SO ₄ to a pH to 2? | | | | CCOD 01 | |
| 2. Upon receipt in the laboratory, is the sample pH measurant recorded to verify that it is preserved? | ured | | | ☐ CCOD 02 | |
| 3. Is the Dichromate reflux method used? | | | _ | | |
| a. Is the ferrous ammonium sulfate titrant standardize daily against primary standard grade K2Cr2O7? | ed | \Box | \Box | ☐ CCOD 03 | |
| b. Is 0.025 N K ₂ Cr ₂ O ₇ used for samples below 50 mg | 3/1? | | | CCOD 04 | |
| c. Is a blank run with each set of samples? | | | | CCOD 05 | |
| d. Is at least 0.5 ml of titrant used in the titration of t excess dichromate for the majority of samples? | the | \Box | | CCOD 06 | |
| e. Is HgSO4 used to complex chloride? | | | | CCOD 07 | |
| s the automated colorimetric method used? | | | | ☐ ccop 68 | AR304112 |
| s the manual colorimetric method used? a. Are digestion tubes heated in a block heater | | لـا | | | |
| or oven at 150°C for 2 hrs.? | | | | | |
| b. Is absorbance read @ 600 nm in a spectrophotomer | ter? | | | | |

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ON-SITE LABORATORY EVALUATION

| | NA. | YES | <u>NO</u> | COMMENTS |
|--|-----|------|-------------|----------|
| HARDNESS, TOTAL (Holding 6 mos.) | | | | |
| Are samples preserved with acid (HNO₃ or H₂SO₄) to pH<2? Is the EDTA titrimetric method used? | | | CHRD 01 . | |
| a. Is the EDTA titrant standardized against CaCO ₃ and labeled properly? | | | | |
| b. Is the EDTA titrant approximately 0.01M?3. Is the automated colorimetric (calmagite) method used? | | | ☐ CHRD 03 . | |
| 4. Is the hardness calculated from Ca+Mg values determined by atomic absorption? | | | | |
| HYDROGEN ION (pH) (Analyze immediately) | | | | |
| Is an electronic pH meter with temperature compensation used? | | | ☐ CpH 01 | |
| 2. Are electrodes stored according to the manufacturer's recommendations? | | Ø | ☐ CpH 02 | |
| 3. Are the electrodes filled with sufficient quantity of electrolyte? | | 四 | ☐ CpH 03 | |
| CONDUCTIVITY (Holding 28 Days) | | | | |
| Are samples measured at 25^oC or is a temperature correction made? | | | CCON 01 | |
| 2. Has the cell constant of the conductance cell been determined and permanently recorded? | | abla | CCON 02 | |
| METHYLENE BLUE ACTIVE SUBSTANCES (Holding 48 Hrs.) | | | | |
| 1. Is MBAS being determined by the methylene blue method? | | | ☐ CMBA 01 | |
| 2. Is LAS reference material available and used in the preparation of standards? | | | ☐ CMBA 02 | |
| 3. Is the determination of absorbance done at 652 nm against a blank of chloroform? | | | ☐ CMBA 03 | |
| TURBIDITY (Holding 48 Hrs.) | | | | |
| Is the nephelometric method used? a. Are samples with turbidity greater than 40 NTU diluted | | | | |
| with turbidity-free water? b. Are sample tubes clear, colorless glass which are clean | | | CTUR 01 | |
| and have no scratches? | | | CTUR 02 | |
| COLOR (Holding 48 Hrs.) | | | | |
| Is the visual comparison method used? a. Is interference due to turbidity removed by filtration | | | | |
| or centrifugation? b. Is the pH of the sample measured and reported with | | | CCOL 01 | |
| the result? | | | | |
| c. Are platinum-cobalt standards used?d. Are color disc standards calibrated against platinum- | | | | |
| cobalt standards every 6 months? 2. Is the spectrophotometric method used? | | | CCOL 04 | |
| 3. Is the ADMI method used? | | | | ANJU41 |

ON-SITE LABORATORY EVALUATION

| | <u>NA</u> | <u>YES</u> | <u>NO</u> | COMMENTS |
|---|-------------|------------|------------|--|
| RESIDUE, (T.D.S.), (TOTAL FILTERABLE RESIDUE) | | | | - |
| (Holding 48 Hrs.) | | | | |
| Does the desiccator have suitable dessicant and indicator? | | | CTDS 01 | |
| 2. Is an analytical balance capable of weighing to 0.1 mg available? | | | CTDS 02 | |
| 3. Are glass fiber filter discs used? | | | CTDS 03 | |
| 4. Are samples for total dissolved solids dried at 180°C? | | | CTDS 04 | |
| 5. a. Does the dissolved residue, when weighed, yield <200 mg? | | | CTDS 05 | |
| b. If not, is smaller aliquot used? | | | CTDS 06 | |
| (Holding 7 Days) | | | | |
| 1. Is the residue dried at 103-105°C? | | | CTSS 01 | |
| RESIDUE, TOTAL SOLIDS (Holding 7 Days) | = | | | |
| 1. Is sample dried at 103-105°C until weight is constant? | | | | |
| CHLORIDE (No Pres., Holding 28 Days) | | | | |
| 1. Is the argentometric (silver nitrate) method used? | | | | |
| a. Is the AgNO3 titrant standardized against NaC1 dried at | | _ | | |
| 140°C and labeled properly? | ليا | | CCLD 01 | |
| b. Is interference due to sulfide, sulfite or thiosulfate removed with H ₂ O ₂ ? | | | CCLD 02 | |
| 2. Is the mercuric nitrate method used? | | | | |
| a. Is the pH adjusted to 2.5? | | | CCLD 03 | |
| b. Is a 1 or 5 ml microburet used for titration? | 1000 | | CCLD 04 | |
| c. Is the NaCL standard dried at 600°C for 1 hour? | 片 | B | CCLD 05 | 325.2 |
| 3. Is the automated ferricyanide method used?4. Is the ion chromatographic method used for drinking water? | | | | |
| FLUORIDE (No Pres., Holding 28 Days) | | | | |
| 1. Are water samples distilled? | | | CFLR 01 | |
| 2. Is the specific ion electrode method used? | | | | |
| a. Are both samples and standards analyzed at room | | | CFLR 02 | |
| temperature? 3. Is the SPADNS method used? | ب | لــا | LI CPER UZ | ************************************** |
| a. Is the SPADNS solution stored in an amber bottle and | | | | |
| protected from direct sunlight? | | | CFLR 03 | |
| b. Is sodium arsenite used to remove residual chlorine? | | | CFLR 04 | |
| 4. Is the automated complexone method used? | | | | .4 |
| a. Is the working color reagent prepared fresh every 3 or 4 days? | \Box | থে | CFLR 05 | 3403 |
| CHLORINE RESIDUAL (No Pres., No Holding) | | | | |
| 1. a. Is chlorine residual determined by iodometric titration, | | | | |
| DPD colorimetric or DPD titrimetric methods? | | | CCLR 01 | |
| b. In the iodometric titration is the excess reducing agent | | | | |
| back-titrated with iodine or iodate solutions? | | | CCLR 02 | |
| c. In the DPD colorimetric method are kits with color | | _ | | AR304114 |
| wheels, and reagent packets used? | لا | لا | u , | - F 5 5 |
| d. Is the chlorine residual determined by specific ion electrode? | | | | |
| e. Is the starch end-point method used? | | | | |

ON-SITE LABORATORY EVALUATION

| Δħ | AMONIA (Holding 28 Days) | <u>NA</u> | <u>YES</u> | <u>NO</u> | | COMMENTS |
|----|--|---------------|-------------|---------------|-----------|---------------|
| | | | | | | |
| | Are samples preserved with H2SO4 to pH 2 at time of collection? | | | | CAMM 01 | |
| | Upon receipt in the laboratory, is the pH measured and recorded? | | | | CAMM 02 | |
| | Are samples analyzed within 28 days of collection? | Ш | لــا | لــا | CAMM 03 | |
| 4. | Is a manual distillation at pH 9.5 used? | | | $\overline{}$ | | |
| | a. Do you use macro or micro distillation equipment? | ш | | لا | CAMM 04 | |
| | b. Are stills steamed with ammonia-free water prior | | | | | |
| | to distillation of samples and the distillate checked | | | | CAMM 05 | |
| | for residual NH3? | | | | CAMIN 05 | |
| | c. Is chlorine residual removed by thiosulfate or arsenite prior to distillation? | | | | CAMM 06 | |
| _ | Is Nesslerization method used following distillation | _ | | _ | CAIMIN OU | |
| Э. | (for 0.05 to 1.0 MGNH ₃ -N/L)? | | | | | |
| | a. Is 2 ml of Nessler reagent added to raise the | | | | | |
| | alkalinity to the desired level? | | | | CAMM 07 | |
| | b. Is the same contact time used for samples | | | | | |
| | standards and blanks? | | | | CAMM 08 | |
| | c. Is a 30 min. contact time allowed for low | | | | | |
| | concentration samples? | | | | CAMM 09 | |
| 6. | Is the selective ion method used | | | | | |
| | (for 0.05 to 1.0 mgNH ₃ ·N/L)? | | | _ | | |
| | a. Is the pH of the sample maintained at greater than 11? | 닏 | 닏 | 닐 | | |
| | b. Is NaOH added to samples prior to electrode immersion? | 닏 | ليا | 닏 | CAMM 11 | |
| | c. Are low concentration standards run first? | Ш | Ш | | CAMM 12 | |
| 7. | Is the automated phenate method used? | | | | | |
| | a. If HgCl2 is used as a preservative, is an equivalent | $\overline{}$ | | _ | | |
| | amount added to NH ₃ standards? | <u></u> | لسا | لــا | CAMM 13 | |
| | b. If H ₂ SO ₄ is used as a preservative, is H ₂ SO ₄ added | $\overline{}$ | | | CAMMA 14 | |
| _ | to wash water and standards? | \vdash | H | H | | |
| 8. | Is titration method used (for 0.05 to 1.0 MGNH3-N/L. | | H | H | | |
| | a. Is H₂SO₄ 0.02N?b. Is a blank carried through all the steps of the procedure? | H | H | 一百 | | |
| | | | | _ | | |
| NI | TRATE | ٠. | | | | |
| 1. | Are drinking water samples analyzed within 24 hours of | | _ | | | • |
| | collection? | \Box | | لـا | CNAT 01 | |
| 2. | Are wastewater samples analyzed within 48 hours of | 区 | | ٠ | | |
| | collection? | | | , ^ [| CNAT 02 | |
| 3. | If not, are samples preserved with H ₂ SO ₄ to pH 2 at | | Ø | | 0NAT 00 | |
| 4 | time of collection for NO ₃ /NO ₂ ? | ب | سي | ч | CNAT 03 | |
| 4. | Is the brucine method used? a. Are samples filtered if turbid? | | \Box | | CNAT 04 | |
| | b. Is the temperature of the waterbath 95 · 100°C? | 一 | | | | |
| | c. Is the stock nitrate STD 100 mg/l, preserved with | | | | 0.17.1.00 | |
| | chloroform and kept no longer than 6 months? | | | | CNAT 06 | |
| | d. Is the brucine-sulfanilic acid reagent stored at 4°C | | | | | |
| | in a dark bottle? | | | | CNAT 07 | |
| | e. Is residual chlorine removed by adding sodium | | | | | |
| | arsenite solution (1 drop/0.1 mg/l)? | | | | CNAT 08 | |
| 5. | Is the manual cadmium reduction method used? | - | _ | | • | |
| | a. Is interference due to turbidity removed? | | | لسا | CNAT 09 | |
| | b. Is a nitrate and nitrite standard passed through the | | | , | 1 | |
| | column with each run to check recovery? | 닏 | | <u> </u> | CNAT 10 | 6 000. |
| _ | c. Is the column reactivated when the value of F>0.33? | ليا | ليا | L | CNAT 11 | |
| 6. | Is the automated cadmium reduction method used? | | | | `- | |
| | a. Is a nitrate and nitrite standard run with each batch of | | ത് | _ | CNAT 12 | 35 3 Z. |
| 7 | samples to check column efficiency? Is the automated hydrazine reduction method used? | H | | | CNAT 13 | |
| | Is the automated hydrazine reduction method used: | | Ī | | | |
| ٠. | The same and a manage appropriate the same against the same manage transfer. | | | | | |

ON-SITE LABORATORY EVALUATION

| NITRITE | <u>NA</u> | <u>YES</u> | <u>NO</u> | COMMENTS |
|---|-----------|------------|-------------------------------------|--|
| Are samples cooled to 4°C and anlyzed within 48 hrs. of collection if not preserved? Is the Diazotization method used? a. Is the nitrite stock solution standardized against standard permanganate and labeled properly? |] [E] | | CNIT 01 | |
| b. Are turbid samples filtered through a 0.45 micron filter? | | d | CNIT 03 | |
| KJELDAHL NITROGEN, TOTAL | | | | |
| Are samples preserved with H₂SO₄ to pH 2 and analyzed within 28 days of collection? Is the 0.020 N H₂SO₄ standardized against Na₂CO₃ and | | | CTKN 01 | |
| properly labeled? 3. Is the distillate from the digestion collected below the | | | CTKN 02 | |
| surface of the boric acid? | | | CTKN 03 | |
| ORTHOPHOSPHATE (Pres. Filter Immed.) | - | | | |
| Are samples cooled to 4°C and analyzed within 48 hrs. of collection? Is the ascorbic acid method used? | | | CORP 01 | |
| a. Is the ammonium molybdate solution stored in plastic at 4°C? | | | ☐ CORP 03 | |
| b. Is the 0.1 M. ascorbic acid stored at 4°C and prepared fresh weekly? | | | CORP 04 | |
| c. Is the combined reagent prepared daily with all reagents at room temperature prior to mixing? | | | CORP 05 | |
| PHOSPHORUS, TOTAL (Pres. H ₂ SO ₄ to pH<2) (Holding 28 Days) | | | | |
| Is an acid-persulfate digestion used for wastewater samples? | | | ☐ CTPH 01 | |
| 2. Is the ascorbic acid method used to determine total phosphorus after the digestion? | | | ☐ CTPH 02 | |
| ORGANIC CARBON, TOTAL (Holding 28 Days) | | | | |
| Are samples preserved with H₂ SO₄ or HCl to pH 2 at time of collection? | | ্তা . | □ стос 01 | |
| Upon receipt in the laboratory, is the pH measured and recorded? | | <u> </u> | CTOC 02 | the state of the s |
| 3. Is the combustion-infrared method used? a. Is inorganic carbon removed by decomposition with acid or alternatively is a correction made for the inorganic fraction? b. Is a methane detection technique used in place of IR? 4. Is analysis performed within 28 days? | | 80B | ☐ CTOC 03 ☐ CTOC 04 ☐ CTOC 05 | |
| 5. Is the instrument being calibrated daily with at least 3 standards? | | 回, | CTOC 06 | |
| 6. Have samples been checked with potassium acid phthalate for recovery? | | ø(| CTOC 07 | |
| 7. Is an external reference sample such as E.M.S.L.Q.C. analyzed at least yearly? | | | ☐ CTOC 08 | |
| 8. Are standards prepared at least monthly? SULFATE (Pres Cool to 4°C - Holding 28 Days) | لــا | لسطا | CTOC 09 | |
| 1. Is the gravimetric method used? | | | | |
| a. Is silica removed by treatment with HC1 and filtering? b. Is the barium sulfate precipitate washed with distilled | | | CSFA 01 | ************************************** |
| water to remove chlorides? c. Is the residue ignited at 800°C? 2. Is the turbidimetric method used? | | | ☐ CSFA 02 ☐ CSFA 03 | AR304116 |
| a. Are the samples stirred for exactly 1 minute after the addition of BaCl ₂ ? | | | CSFA 04 | |

ON-SITE LABORATORY PROCEDURES

| | <u>.VA</u> | <u>YES</u> | <u>NO</u> | <u>COMMENTS</u> |
|---|------------|------------|--|-----------------|
| SULFATE — Continue 2. b. Are both sample. d standards read at 4 minutes after stirring? c. Are blanks used to correct for color or turbidity? 3. Is the automated chloranilate method used? | . = | | CSFA 05 | |
| a. Are interferences due to Ca, Al, and Fe removed by an ion exchange column? 4. Is the ion chromatography method used for drinking water? | | | CSFA 07 | |
| <u>SULFIDE</u> (Pres 4 ⁰ C,Zn Acetate + NaOH to pH>9 · · · · · · · · · · · · · · · · · · · | | | | |
| Is the Methylene Blue method used? a. Is the methylene blue solution standardized against a known solution and adjusted so that 1 drop = 1.0 mg/l sulfide? b. Is the titrimetric (Iodine) method used? | ` | | CSFD 01 | |
| SULFITE (No Preservation) | | | | |
| Is the titrimetric iodine-iodate method used? Are samples analyzed on site? | | | CSFT 01 | |
| CYANIDE | | / | / | |
| 1. Are samples analyzed within 14 days of collection? | | \Box | CCYN 01 | |
| Are samples preserved with NaOH to pH 12 + 0.6 G ascorbic acid? | | <u></u> | CCYN 02 | |
| 3. Upon receipt in the laboratory, is the pH measured and recorded?4. If chlorinated, do you remove sulfide as Cd sulfide? | | | CCYN 03 | 4-11 |
| 5. Is the titrimetric method used? 6. Is a manual distillation with MgCl₂ done? a. Is the AgNO₃ standardized against NaCl and labeled properly? b. Is a blank run with each set of samples? 7. Is the colorimetric method used? | | | CCYN 05 | |
| a. Is Chloramine T prepared weekly and stored in refrigerator? | | <u>v</u> | CCYN 07 | |
| b. Is the stock cyanide solution standardized weekly against AgNO ₃ ? | | | ☑ CCYN 08 | |
| OIL AND GREASE (Holding 28 Days) | | | | |
| Are samples collected in glass containers? Are samples preserved with H₂SO₄ to pH<2? Is a liquid-liquid extraction with freon used? Is the oil and grease content determined gravimetrically? | | | ☐ CONG 01 ☐ CONG 02 ☐ CONG 03 ☐ CONG 04 | |
| PHENOLS (Holding 28 Days) | | | | |
| Are samples collected in glass containers? Are samples preserved with 1 g. CuSO₄ and H₃PO₄ to pH<2? | | | ☐ CPHN 01 ☐ CPHN 02 | |
| 3. Upon receipt in the laboratory, is the pH measured and recorded? 4. Are samples analyzed within the pH measured and recorded? 5. Is the colorimetric 42 P method with distillation used? | | | ☐ CPHN 03 ☐ CPHN 04 ☐ CPHN 05 | without the in |
| 6. Is the colorimetric 4AAP method for halogenated phenols used? | | | ☐ CPHN 06 | AR304117 |

ON-SITE LABORATORY PROCEDURES

| | <u>.V.4</u> | YES | <u>.vo</u> | COMMENTS |
|--|----------------|---------|------------------------|-----------|
| ALUMINUM | | | | |
| 1. Is the Errochrome Cyanine R method used? a. Is an EDTA complexed aliquot run as a blank? b. Is Fe and Mn. orer ference removed with Ascorbic acid? | | = | CALU 01 CALU 02 | |
| c. Is F compensated for by addition of F to standards? | _ _ | | CALU 03 | |
| d. Are interferences due to polyphosphates and alkalinity remulard by treatment with H2SO4? | 二 | _ | CALU 04 | |
| ARSENIC | | | | |
| 1. Is the silver diethyldithocarbamate method used? | \Box | \Box | | |
| a. Are samples preserved with H₂SO₄ to avoid negative interference from HNO₃? b. Is a lead acetate scrubber used? | | | ☐ CARS 01 ☐ CARS 02 | |
| BERYLLIUM | | | | |
| 1. Is the Aluminon method used? | | | | |
| a. Is EDTA added to eliminate AI, Co, Cu, Fe, Mn, Ni, Ti, Zn. and Zr interferences? | | | CBER 01 | |
| BORON | | | | • |
| 1. Is the Curcumin method used? | | | | |
| a. Is water bath maintained at 55± 2° C? b. Is Curcum is prepared fresh weekly and refrigerated? | | .][| CBOR 01 | |
| c. Is interference due to the hardness removed by ion exchange of filtering of final sample? | | | CBOR 03 | |
| d. Is the strying time carefully controlled for both samples and standa.ds? | | | CBOR 04 | · |
| CADMIUM | | | | |
| - Company of the Comp | | | | |
| Is the Dithizone method used? a. Are pastewater samples digested with HNO₃ + H₂SO₄ | اسما | | | |
| or HNO ₁ - HC10 ₄ ? | | | CCAD 01 | |
| b. Is chloroform supplied in containers with metal- lined cups redistilled? | | | CCAD 02 | |
| c. During the final extraction, is the room darkened or amber glassware gsed? | | | CCAD 03 | |
| CALCIUM | | | | |
| 1. Is the EDTA titrimetric method used? | | | CCAL 01 | |
| a. Is the EDTA titrant standardized against CaCO ₃ ? b. Is EDTA titrant approximately 0.01 M and labeled | - | <u></u> | CCAL UI | |
| property? | | | CCAL 02 | - |
| c. After the pH is raised to 12-13, is the sample titrated immediately? | | . 🗆 | CCAL 03 | |
| CHROMIUM | | | | |
| 1. Is the Dipnenylcarbazide method used? | | | | |
| a. Is KMnO₁ used to oxidize CrIII to CrVI? b. Is permanganate interference removed by reduction | | | CCHR 01 | |
| with azios? | | | CCHR 02 | AR201.110 |
| c. Are standards processed in the same manner as samples? | | | CCHR 03 | 411204118 |

ON-SITE LABORATORY PROCEDURES

| CHROMIUM VI (Pres Cool 4°C - Holding 24 Hrs.) | .VA | YES | NO | COMMENTS |
|--|-----|-----|-------------------------|----------|
| Is the analysis performed in the field? Is the sample screened for Cr⁺⁶ using a total chromium determination? | | | | |
| COPPER | | | | |
| Is the neocuproine method used? a. Is glass redistilled or deionized distilled water used? | | | CCOP 01 | |
| b. Are samples digested with H₂SO₄ and HNO₃ to remove cyanide and sulfide interference? c. Is the blank treated in the same manner as the sample? d. Is the bicinchoninate method used? | | | CCOP 02 CCOP 03 | |
| IRON | | | | |
| Is the phenanthroline method used? a. Are reagents stored in glass stoppered bottles? b. Are working-standard iron solutions prepared daily? | | | CFER 01 CFER 02 | |
| MANGANESE | | | - | |
| Is the persulfate method used? a. Is interference from NaCl removed by addition of mercuric sulfate? | | | ☐ CMAN 01 - | |
| b. Is the manganese standard aged in sunlight or heated, then standardized against sodium oxalate?2. Is the periodate method used? | | | CMAN 02 | |
| a. Are reducing agents removed or destroyed before the periodate oxidation? b. Is phosphoric acid added to complex ferric iron? c. Are corrections for turbidity or interfering color made? | | | CMAN 03 CMAN 04 CMAN 05 | |
| SODIUM | | | | |
| Is the flame photometric method used? Is particulate matter removed by filtration? Are all solutions stored in plastic bottles? | | | CSOD 01 CSOD 02 | |
| SILVER | | | | |
| Is the Dithizone method used? a. Is the stock dithizone solution extracted with CCI4 to | | | | |
| remove Cu, then stored in the dark or in an amber bottle? b. Is all glassware washed with chromic acid and | | | CSIL 01 | |
| 1 + 1 HNO t then treated with a silicone coating? c. Are urea solutions discarded when a red film develops? | | | CSIL 02 | |
| VANADIUM | | | | |
| Is the Gallic Acid method used? a. Is there a water bath capable of 25 ± 0.5 °C available? b. Is the absorbance measured exactly 60 min. after | | | CVAN 01 | |
| the addition of gallic acid? c. Are interferences due to Cu and Fe eliminated by dilution? | | | CVAN 02 CVAN 03 | |
| ZINC | | | | · · · |
| 1. Is the dithizone method used? | | | | |
| a. Is the NH4 OH solution made from NH3 or by redistilling NH4OH? b. Is the dilute sodium sulfide solution prepared fresh | | | CZIN 01 | |
| just before use? c. Are blanks reproducible? | | | CZIN 03 | AR304119 |

ON-SITE LABORATORY EVALUATION

ATOMIC ABSORPTION AND METAL PROCEDURES

| | | <u>NA</u> | <u>YES</u> | <u>.vo</u> | COMMENTS |
|-------------------|--|---------------------|------------|------------|---|
| 2. 3. 4. | Does the instrument have the following: Background Correction - Continuum Source Stripchart Recorder Double Beam Graphite Furnace Auto Sampler | | য়েবাবাবাব | | |
| | Are trace metals samples preserved with HNO3 to pH of 2 at time of collection? (Holding: Hg - 28 days, others 6 mos.) | \Box | ☐´ | \Box | CAAS 01 |
| 7. | Are samples for dissolved metals filtered through 0.45 u membrane filter and the filtrate preserved with HNO3? | | V | | CAAS 02 |
| | Is an acid digestion done on total metals samples for drinking water? | | Σ. | | |
| 10. 11. 12. | Upon receipt in laboratory is sample pH measured & recorded? Is deionized-distilled or double-distilled water used? Is glassware acid washed? Are all the required lamps available for parameters requested? List lamps available and underline multi-element lamps. | | RINDIST | | CAAS 03 |
| | Are all the following fuel mixtures available? Air-Acetylene Nitrous oxide-acetylene Argon-hydrogen (Hydride Generation · Zn+SnCl2) Circle if not available | | | , | CAAC 07 |
| 15. | In the graphite furnace method, is each sample matrix examined for interference effects by the method of standard additions? | | | | CAAM 08 , |
| 16. | Are 10% check standards run for the graphite furnace method to monitor when the furnace should be changed? | | 过 | | |
| 7. | For Cd, Cr, Pb and Zn are low level samples extracted into MIBK after chelation of the desired metal with APDC or is CHCl3 used as a solvent for PDCA extraction? | | | | CAAM 09 |
| 18. | In the determination of low level chromium, is Cr III oxidized to Cr VI prior to extraction? | $ \mathbf{\nabla} $ | . 🗆 | | CAAM 10 |
| 19. | Are Se and As converted to the gaseous hydride with SNCI2 + Zn metal and determined in an argon-hydrogen flame? | ರ [′] | | | CAAM 11 |
| 20. | Is a nitrous oxide flame used for AI, Ba, Be, Mo, Ti, Sn & V? | \Box / | | | CAAM 12 |
| 21. | If Ba is determined using an air-acetylene flame, is La added to both samples and stds? | Ź | | | CAAM 13 |
| 22. | For Al, Ba, Na and Ti analysis, is K added to both samples and standards to eliminate ionization of the measured species? | | | | CAAM 14 |
| 23. | For Ca and Mg analyses, is La added to both samples and standards to eliminate interference? | | | | CAAM 15 |
| 24. | For Cr by graphite furnace and Mn and Fe by direct aspiration analyses, is Ca added to both samples and standards to eliminate interference? | | | · 🗆 | CAAM 16 Terri Lie 17 |
| 25. | Is apparatus available for the determination of Hg by the cold vapor technique of Hatch and Ott? | | Ø | | CAAM 17 |
| 26. | Is a KMnO4 trap, some type of scrubber or venting up the hood used in the apparatus for flameless Hg determination? | | Ó | | CAAM 18 |
| 27. | If only dissolved mercury is to be determined, is the sample filtered through an all glass apparatus before the acid is added? | | \Box | | CAAM 19 |
| 29. 30. 31. | Is persulfate added when determining total Hg? Is KMnO4 added until dark color persists? Are the samples heated for 2 hours at 95°C in a water bath? Is mercury reduced with SnCl2 or SnSO4? For Mo and V analyses is Al added to both samples & standard | | | | CAAM 20 CAAM 21 CAAM 22 CAAM 23 CAAM 24 |

ON-SITE LABORATORY EVALUATION

ICAP AND DCP PROCEDURES

| | | <u>.V.4</u> | $\frac{YES}{NO}$ | - COMMENTS | |
|-----|---|-------------|------------------|------------|-------------|
| 1. | Does the instrument have background correction? | | | | |
| 2. | If DCP, does the instrument have a 3 electrode system, not a 2 electrode system? | | | | |
| 3. | Does the instrument have computer control? | | | | |
| 4. | Is a peristaltic pump used with the system? | | | | |
| 5. | If DCP, are enhancers used? | | | | |
| 6. | Does the instrument have temperature control or is the environment temperature and humidity controlled? | | | | |
| 7. | Are the acids used trace metal grade? | | | | |
| 8. | Is an instrument check standard run 10% of time to check for impurities and spectral interferences? | | | | |
| 9. | Are 10x Instrument Detection Limit spikes run (1 every 20)? | | | | |
| 10. | Is sample digestion documented? | | | | |
| 11. | Is instrument monitored weekly for stability? | | ष्टं 🗆 | | |
| 12. | Is a profile check run every 4 hours if not documented or at least once a shift and documented? | | / a () | | |
| 13. | If there is no peristaltic pump used are samples filtered? | I | , 🗆 🗆 | | |
| 14. | is a white light and a dark current check run at least every 3 months? | | / = | | |
| 15. | Is the correlation coefficient > 0.9999? | | | | |
| 16. | Is a linear range analysis curve run over the range of interest to check for interferences? | | | | |
| 17. | Do you have welding goggles to look at the plasma? (EPA 79 manual) | I | | | |
| 18. | Are the correct lines being used? | | | | |
| 19. | Do you keep an instrument maintenance log? | | | | |
| 20. | Do you have EPA check samples for interference? | | | | |
| 21. | If the argon is not liquid, how pure is it? | | | | |

AR304121

Form DWR-156 5/85

NEW JERSEY DEPARTMENT OF ENVIRONMENTAL PROTECTION OFFICE OF SCIENCE AND RESEARCH ON-SITE LABORATORY EVALUATION

RECORD-KEEPING AND CALIBRATION PRACTICES

| R | ECORD-KEEPING | NA , | YES | NO_ | | COMMENTS |
|----------|--|-------------|-----------|--------|--|-----------|
| 1. | Is the temperature of all BtO.D. incubators recorded daily? | □ | | | CREC 01 | |
| 2. | Is the temperature of all drying ovens recorded daily? | | \square | | CREC 02 | |
| 3. | Is the temperature of all refrigerators recorded daily? | | Œ | | CREC 03 | |
| 4. | Are the laboratory thermometers calibrated against an NBS traceable thermometer and documented? | | ত্র | | CREC 04 | |
| 5. | Is the analytical balance checked monthly with two class S weights, one in the mg range, and one in the gram range, and the data recorded? | | | | CREC 05 | |
| 6. | Is a record available of yearly service on the analytical balance? | | 四二 | | CREC 06 | |
| 7. | Is the pH meter checked daily, or before use, by setting the meter to pH7 then measuring and recording pH's approximately 4 and 10? | | 4 | | CREC 07 | |
| 8. | Is the conductivity of the distilled water supply (satisfactory is conductivity of 2.0-0.5 umho/cm. at 25°C.) checked daily and the data recorded? | | \Box | . 🗆 | CREC 08 | |
| 9. | Is the conductivity meter calibrated daily against a 0.001 M KCI solution and the data recorded? | □, | ପ୍ର | | CREC 09 | |
| 10. | Is the turbidimeter calibrated daily, or before use, with a 40 NTU formazin standard and the data recorded? | | | | CREC 10 | |
| 11. | Is the DO meter calibrated weekly against the Winkler method and the data recorded? | | | | CREC 11 | |
| <u>C</u> | ALIBRATION PRACTICES | | | | | |
|) | Regarding calibration curves, are the following practices in use a. Graph is labeled with parameter, date of calibration and the axes are properly identified as to absorbance or percent transmission and concentration units. b. Computer read-out for regression analysis lists parameter, date of calibration, equation of curve and correlation co-efficient. c. Results reported are within the range of the highest and lowest standard. | | B B | | CCAL 01 CCAL 02 CCAL 03 | |
| 2. | Regarding manual spectrophotometric calibration curves, are the following practices in use? a. A minimum of 5 standards and a blank, with 3 measurements at each point are used to generate the curve. b. A new curve is generated every 6 months. c. The working curve is checked daily or with each run by alternating a low and a high standard and the data are recorded. | | | | CCAL 04 CCAL 05 | |
| 3. | Regarding calibration curves for auto-analyzer analyses, are | | | | | |
| | the following practices in use? a. The baseline is set using appropriate reagents and distilled water and is checked at the end of the run. b. A minimum of 5 standards are used to generate the curve. c. A new curve is generated for each run. d. A marking standard is included with every 20 samples. e. The calibration curve is checked at the end of each run with a low and a high standard and the data are recorded. | 0000 0 | व वव्यव | 0 0000 | CCAL 07 CCAL 08 CCAL 09 CCAL 10 | |
| 4. | Regarding atomic absorption calibration curves, are the following practices in use? | | | , | | 8000t too |
| | a. Working standards are prepared fresh with each run. b. A minimum of 4 standards and a blank are used to | | | | CCÁL 12 | AR304122 |
| | generate a curve. c. A new curve is generated for each run. | | | | CCAL 13 CCAL 14 | , |

ON-SITE LABORATORY EVALUATION

QUALITY CONTROL AND DATA HANDLING .

| | <u>NA</u> | <u>YES</u> | <u>NO</u> | COMMENTS |
|--|-----------|----------------------|---|----------|
| QUALITY CONTROL | | | | |
| Regarding standard solutions, are the following practices in use a. A notebook record is available describing the preparation and standardization of stock standard solutions. b. Are purchased standards checked before use? c. Stock standard solutions and working standards are labeled with reagent, concentration, date prepared and initialed. d. ACS grade or analytical reagent grade chemicals dated when received, are used in the preparation of standard solutions. | | | COCS 01 COCS 02 COCS 03 | notali |
| Regarding the monitoring of precision, are the following practices in use? a. Approximately 1 synthetic known control sample is included with every 20 analyses, and the data presented on an X bar control chart. b. Approximately 1 duplicate of a natural sample is included with every 20 analyses, and the data presented on an R bar range control chart. | ed | | □ CQCP 02 | |
| 3. Regarding the determination of chemical recovery, are the following practices in use? a. A tabulation and control chart are available for recovery data obtained from spiked natural samples (1 for every 20 analyses). 4. Is there an in-house quality control manual outlining Q.C. practices? | | a | ☐ CQCP 03 | |
| DATA HANDLING | | | | |
| Regarding sampling procedures and data handling, data reporting and data retrieval procedures, are the following practices in use? a. Are sample collectors supplied with properly labeled containers, preservatives and sampling instructions? (Get copy of instructions). b. Is there a lab daily work sheet listing sample number, date, time, location, preservation, analyses requested, field measurements by sampler, sampler's initials, date | | ⊠í | ☐ CDAT 01 | |
| and hour received by lab, analysis, date and hour of analysis, analyst's initials? c. Is there a bound lab notebook for recording raw data, calculations, or other notes. d. Is raw data kept for 5 years? e. Is enforcement data kept for 5 years? f. Is there an in-house methods manual available to all analysts? g. Is there a record of chain of custody? h. Is there a chain of custody procedure? | 0 000 000 | विवृद्धि विवृद्धि वि | CDAT 02 CDAT 03 CDAT 04 CDAT 05 CDAT 06 CDAT 07 CDAT 08 | AR3U4123 |